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ENTER TYPE OF SEARCH (SSS), CSS, FAMILY, OR EXACT:sss
ENTER SCOPE OF SEARCH (SAMPLE), FULL, RANGE, OR SUBSET:subset
ENTER SUBSET L# OR (END):l3
ENTER SUBSET SEARCH SCOPE - SAMPLE, FULL, RANGE, OR (END):ful
FULL SUBSET SEARCH INITIATED 08:07:45 FILE 'REGISTRY'
FULL SUBSET SCREEN SEARCH COMPLETED -       310 TO ITERATE

100.0% PROCESSED      310 ITERATIONS                          310 ANSWERS
SEARCH TIME: 00.00.01

L5          310 SEA SUB=L3 SSS FUL L4

=> s l3 not l5
L6          631 L3 NOT L5

=> fil caplus
COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                       ENTRY    SESSION
FULL ESTIMATED COST                                   214.55     214.76

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FILE LAST UPDATED: 11 Jun 2007  (20070611/ED)

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=> s l5
L7          11 L5

=> d bib 1-11

L7   ANSWER 1 OF 11  CAPLUS  COPYRIGHT 2007 ACS on STN
AN   2007:282077  CAPLUS
DN   146:337878
TI   Pyrazolecarboxamide derivatives, process for preparing them, their use as
     antagonists or inverse agonists of cannabinoid CB1 and opioid μ
     receptors
IN   Jagerovic, Nadine; Fernandez Fernandez, Cristina; Goya Laza, Maria Pilar;
     Callado Hernando, Luis Felipe; Meana Martinez, Jose Javier
PA   Consejo Superior de Investigaciones Cientificas, Spain; Universidad del
     Pais Vasco
SO   PCT Int. Appl., 57pp.
     CODEN: PIXXD2
DT   Patent
LA   Spanish
FAN.CNT 1
```

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2007028849	A1	20070315	WO 2006-ES70132	20060907

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

PRAI ES 2005-2196 A 20050908

OS MARPAT 146:337878

RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 2 OF 11 CAPLUS COPYRIGHT 2007 ACS on STN

AN 2006:1005694 CAPLUS

DN 145:377208

TI Preparation of N-substituted-N-(4-piperidinyl)amide derivatives as analgesics

IN Takahashi, Toshihiro; Endo, Tsuyoshi; Shiota, Katsutoshi; Sakuma, Syogo; Yamakawa, Tomio; Shika, Kiichi; Kawasaki, Toru; Imai, Toshiyasu; Hirate, Kenji

PA Nippon Chemiphar Co., Ltd., Japan

SO PCT Int. Appl., 101pp.
CODEN: PIXXD2

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2006101245	A1	20060928	WO 2006-JP306381	20060322

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

PRAI JP 2005-83653 A 20050323

OS MARPAT 145:377208

RE.CNT 21 THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 3 OF 11 CAPLUS COPYRIGHT 2007 ACS on STN

AN 2006:272555 CAPLUS

DN 144:331267

TI Preparation of N-phenyl-N-(4-piperidinyl)amide derivatives as μ opioid receptor antagonists for the treatment of pain

IN Takahashi, Toshihiro; Endo, Tsuyoshi; Shiota, Katsutoshi; Kobayashi, Kunio; Yamakawa, Tomio; Shika, Kiichi; Kawasaki, Toru; Imai, Toshiyasu; Hirate, Kenji

PA Nippon Chemiphar Co., Ltd., Japan

SO PCT Int. Appl., 96 pp.

CODEN: PIXXD2
DT Patent
LA Japanese
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2006030931	A1	20060323	WO 2005-JP17217	20050913

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

PRAI JP 2004-267238 A 20040914
OS MARPAT 144:331267
RE.CNT 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 4 OF 11 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2005:346988 CAPLUS
DN 142:392299
TI Preparation of aniline- and aminopyridine-derivatives as 5-HT1F receptor agonists
IN Blanco-Pillado, Maria-Jesus; Cohen, Michael Philip; Filla, Sandra Ann; Hudziak, Kevin John; Kohlman, Daniel Timothy; Benesh, Dana Rae; Victor, Frantz; Xu, Yao-Chang; Ying, Bai-Ping; Zacherl, Deanna Piatt; Zhang, Deyi
PA Eli Lilly and Company, USA
SO PCT Int. Appl., 127 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2005035499	A1	20050421	WO 2004-US25607	20040903

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

AU 2004280320	A1	20050421	AU 2004-280320	20040903
CA 2537936	A1	20050421	CA 2004-2537936	20040903
EP 1663971	A1	20060607	EP 2004-780442	20040903

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, HR

CN 1849307	A	20061018	CN 2004-80026400	20040903
BR 2004014241	A	20061107	BR 2004-14241	20040903
JP 2007505105	T	20070308	JP 2006-526084	20040903
US 2006287363	A1	20061221	US 2006-569109	20060221
IN 2006KN00450	A	20070202	IN 2006-KN450	20060227
NO 2006001584	A	20060606	NO 2006-1584	20060407
PRAI US 2003-502780P	P	20030912		

WO 2004-US25607 W 20040903
OS MARPAT 142:392299
RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 5 OF 11 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2003:855655 CAPLUS
DN 139:350636
TI Preparation of amino heteroaryl amides for use in pharmaceutical compositions for the treatment of angiogenesis mediated diseases such as cancer
IN Patel, Vinod F.; Askew, Benny; Booker, Shon; Chen, Guoqing; Dipietro, Lucian V.; Germain, Julie; Habgood, Gregory J.; Huang, Qi; Kim, Tae-seong; Li, Aiwen; Nishimura, Nobuko; Nomak, Rana; Riahi, Babak; Yuan, Chester Chenguang; Elbaum, Daniel
PA Amgen Inc., USA
SO U.S. Pat. Appl. Publ., 148 pp., Cont.-in-part of U.S. Ser. No. 46,622.
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2003203922	A1	20031030	US 2002-197918	20020717
	US 7102009	B2	20060905		
	US 2003195230	A1	20031016	US 2002-46622	20020110
	US 7105682	B2	20060912		
	CN 1538836	A	20041020	CN 2002-806467	20020111
	ZA 2003005198	A	20040630	ZA 2003-5198	20030704
	CA 2492045	A1	20040122	CA 2003-2492045	20030715
	WO 2004007481	A2	20040122	WO 2003-US22275	20030715
	WO 2004007481	A3	20040219		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	AU 2003263784	A1	20040202	AU 2003-263784	20030715
	EP 1562933	A2	20050817	EP 2003-764755	20030715
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	JP 2006502118	T	20060119	JP 2004-521922	20030715
	US 2006194848	A1	20060831	US 2006-417329	20060502
PRAI	US 2001-261882P	P	20010112		
	US 2001-323808P	P	20010919		
	US 2002-46622	A2	20020110		
	US 2002-197918	A	20020717		
	WO 2003-US22275	W	20030715		

OS MARPAT 139:350636
RE.CNT 84 THERE ARE 84 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 6 OF 11 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2002:676007 CAPLUS
DN 137:216945
TI Preparation of substituted 2-(1H-indazol-6-ylamino)nicotinamides for treating KDR-related diseases
IN Chen, Guoqing; Adams, Jeffrey; Bemis, Jean; Croghan, Michael; Dipietro, Lucian; Dominguez, Celia; Elbaum, Daniel; Germain, Julie; Huang, Qi; Kim,

Joseph L.; Ouyang, Xiaohu; Patel, Vinod F.; Smith, Leon M.; Tasker, Andrew; Xi, Ning; Xu, Shimin; Yuan, Chester Chenguang; Kim, Tae-Seong

PA Amgen Inc., USA

SO PCT Int. Appl., 395 pp.
CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002068406	A2	20020906	WO 2002-US3064	20020111
	WO 2002068406	A3	20030424		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	US 2003195230	A1	20031016	US 2002-46622	20020110
	US 7105682	B2	20060912		
	CA 2434178	A1	20020906	CA 2002-2434178	20020111
	AU 2002253890	A1	20020912	AU 2002-253890	20020111
	HU 200302719	A2	20031128	HU 2003-2719	20020111
	EE 200300325	A	20031215	EE 2003-325	20020111
	JP 2004527499	T	20040909	JP 2002-567920	20020111
	CN 1538836	A	20041020	CN 2002-806467	20020111
	EP 1467721	A2	20041020	EP 2002-723086	20020111
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	ZA 2003005198	A	20040630	ZA 2003-5198	20030704
	BG 108013	A	20040430	BG 2003-108013	20030721
	US 2006194848	A1	20060831	US 2006-417329	20060502
PRAI	US 2001-261882P	P	20010112		
	US 2001-323808P	P	20010919		
	US 2002-46622	A	20020110		
	WO 2002-US3064	W	20020111		

OS MARPAT 137:216945

L7 ANSWER 7 OF 11 CAPLUS COPYRIGHT 2007 ACS on STN

AN 2000:530560 CAPLUS

DN 133:261089

TI Synthesis and evaluation of 4-(N,N-diarylamino)piperidines with high selectivity to the δ-opioid receptor: a combined 3D-QSAR and ligand docking study

AU Podlogar, Brent L.; Poda, Gennady I.; Demeter, David A.; Zhang, Sui-Po; Carson, John R.; Neilson, Lou Anne; Reitz, Allen B.; Ferguson, David M.

CS Department of Chemistry, Bayer Research Center, West Haven, CT, 06516, USA

SO Drug Design and Discovery (2000), 17(1), 34-50
CODEN: DDDIEV; ISSN: 1055-9612

PB Harwood Academic Publishers

DT Journal

LA English

RE.CNT 61 THERE ARE 61 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 8 OF 11 CAPLUS COPYRIGHT 2007 ACS on STN

AN 1999:801468 CAPLUS

DN 132:57145

TI Ink-jet recording materials and ink-jet recording inks

IN Sugiyama, Jun; Ohnishi, Hiroyuki; Sano, Yukari

PA Seiko Epson Corp., Japan
SO Jpn. Kokai Tokkyo Koho, 12 pp.
CODEN: JKXXAF
DT Patent
LA Japanese
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 11348418	A	19991221	JP 1998-264488	19980918
PRAI JP 1998-96214	A	19980408		

L7 ANSWER 9 OF 11 CAPLUS COPYRIGHT 2007 ACS on STN
AN 1999:595124 CAPLUS
DN 131:228549
TI Preparation of (oxalylamino)benzoic acid derivatives and analogs as modulators of protein tyrosine phosphatases (PTPases)
IN Richter, Lutz Stefan; Andersen, Henrik Sune; Vagner, Josef; Jeppesen, Claus Bekker; Moller, Niels Peter Hundahl; Branner, Sven; Su, Jing; Bakir, Farid; Judge, Luke Milburn
PA Novo Nordisk A/S, Den.; Ontogen Corporation
SO PCT Int. Appl., 100 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 6

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 9946236	A1	19990916	WO 1999-DK122	19990311
W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW				
RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
US 6225329	B1	20010501	US 1999-265069	19990309
AU 9927136	A	19990927	AU 1999-27136	19990311
EP 1062199	A1	20001227	EP 1999-907333	19990311
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, PT, IE, FI				
JP 2002506055	T	20020226	JP 2000-535619	19990311
ZA 9902029	A	19990927	ZA 1999-2029	19990312
PRAI DK 1998-342	A	19980312		
DK 1998-345	A	19980312		
DK 1998-472	A	19980403		
DK 1998-479	A	19980403		
DK 1998-940	A	19980715		
US 1998-82913P	P	19980424		
US 1998-82914P	P	19980424		
US 1998-93638P	P	19980721		
WO 1999-DK122	W	19990311		

OS MARPAT 131:228549
RE.CNT 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 10 OF 11 CAPLUS COPYRIGHT 2007 ACS on STN
AN 1995:227441 CAPLUS
DN 122:105695
TI Carbostyril oxytocin receptor antagonists
IN Freidinger, Roger M.; Pawluczyk, Joseph M.; Pettibone, Douglas J.; Williams, Peter D.
PA Merck and Co., Inc., USA
SO U.S., 177 pp.
CODEN: USXXAM

DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 5356904	A	19941018	US 1992-957491	19921007
	WO 9519773	A1	19950727	WO 1994-US847	19940119
	W: CA, JP				
	RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
PRAI	US 1992-957491		19921007		

OS MARPAT 122:105695

L7 ANSWER 11 OF 11 CAPLUS COPYRIGHT 2007 ACS on STN
AN 1991:81619 CAPLUS
DN 114:81619
TI Preparation of carbostyril derivatives as vasopressin antagonists
IN Ogawa, Hidenori; Miyamoto, Hisashi; Kondo, Kazumi; Yamashita, Hiroshi; Nakaya, Kenji; Tominaga, Michiaki; Yabuuchi, Yoichi
PA Otsuka Pharmaceutical Co., Ltd., Japan
SO Eur. Pat. Appl., 364 pp.
CODEN: EPXXDW
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 382185	A2	19900816	EP 1990-102404	19900207
	EP 382185	A3	19910918		
	EP 382185	B1	19940615		
	R: CH, DE, DK, ES, FR, GB, IT, LI, NL, SE				
	ES 2056259	T3	19941001	ES 1990-102404	19900207
	JP 03173870	A	19910729	JP 1990-31360	19900208
	JP 07068218	B	19950726		
	CN 1046529	A	19901031	CN 1990-100657	19900210
	CN 1036394	B	19971112		
	KR 9711153	B1	19970707	KR 1990-1705	19900210
	US 5225402	A	19930706	US 1991-762736	19910918
	US 5436254	A	19950725	US 1993-125667	19931102
	US 5652247	A	19970729	US 1994-359081	19941214
PRAI	JP 1989-31580	A	19890210		
	JP 1989-102699	A	19890421		
	JP 1989-181440	A	19890713		
	JP 1989-232333	A	19890907		
	US 1990-478181	B1	19900209		
	US 1991-762736	A3	19910918		
	US 1992-846941	A1	19920306		

OS MARPAT 114:81619

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> d his

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     FILE 'REGISTRY' ENTERED AT 08:05:15 ON 12 JUN 2007
L1             STRUC
L2          41 S L1
L3         941 S L1 FUL
L4             STRUC
L5         310 SEARCH L4 SSS  SUB=L3 FUL
L6         631 S L3 NOT L5

     FILE 'CAPLUS' ENTERED AT 08:08:14 ON 12 JUN 2007
L7          11 S L5

=> s l6
L8          23 L6

=> d bib 1-23

L8   ANSWER 1 OF 23  CAPLUS  COPYRIGHT 2007 ACS on STN
AN   2006:1012400  CAPLUS
DN   145:383500
TI   Combinations for the treatment of cancer
IN   Chang, David
PA   Amgen Inc, USA
SO   PCT Int. Appl., 48 pp.
     CODEN: PIXXD2
DT   Patent
LA   English
FAN.CNT 1
     PATENT NO.           KIND   DATE        APPLICATION NO.         DATE
     ---------------      ----   --------    --------------------    --------
PI   WO 2006102504         A2    20060928    WO 2006-US10582         20060322
     WO 2006102504         A3    20061207
         W:  AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
             CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
             GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR,
             KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX,
             MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE,
             SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC,
             VN, YU, ZA, ZM, ZW
         RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
             IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ,
             CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH,
             GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA
     US 2006216288         A1    20060928    US 2006-386271          20060321
PRAI US 2005-664381P       P     20050322
     US 2006-386271        T0    20060321
OS   MARPAT 145:383500

L8   ANSWER 2 OF 23  CAPLUS  COPYRIGHT 2007 ACS on STN
AN   2006:342829  CAPLUS
DN   144:390559
TI   Preparation of benzenesulfonamide compounds as N-type calcium channel
     inhibitors
IN   Ohtani, Tazumi; Kambe, Tohru; Kobayashi, Kaoru; Takimizu, Hideyuki; Ito,
     Yoko
PA   Ono Pharmaceutical Co., Ltd., Japan
SO   PCT Int. Appl., 196 pp.
     CODEN: PIXXD2
DT   Patent
LA   Japanese
```

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2006038594	A1	20060413	WO 2005-JP18306	20051003

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

PRAI JP 2004-290916 A 20041004

OS MARPAT 144:390559

RE.CNT 13 THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 3 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN

AN 2006:79149 CAPLUS

DN 144:150365

TI Preparation of ureidopyrazoles as p38 kinase inhibitors

IN De Dios, Alfonso; Li, Tiechao; Martin Cabrejas, Luisa Maria; Pobanz, Mark Andrew; Shih, Chuan; Wang, Yong; Zhong, Boyu; Blas, Jesus Andres; Lopez De Uralde-Garmendia, Beatriz

PA Eli Lilly and Company, USA

SO PCT Int. Appl., 95 pp.
CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2006009741	A1	20060126	WO 2005-US21148	20050615

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

	EP 1609789	A1	20051228	EP 2004-380131	20040623

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, HR

	EP 1761520	A1	20070314	EP 2005-766569	20050615

R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BA

PRAI	EP 2004-380131	A	20040623
	US 2004-592539P	P	20040730
	EP 2004-380174	A	20040823
	US 2004-622492P	P	20041027
	WO 2005-US21148	W	20050615

OS MARPAT 144:150365

RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 4 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2005:1348961 CAPLUS
DN 144:69825
TI Preparation of ureidopyrazoles as p38 kinase inhibitors
IN De Dios, Alfonso; Li, Tiechao; Martin-Cabrejas, Luisa Maria; Pobanz, Mark Andrew; Shih, Chuan; Wang, Yong; Zhong, Boyu
PA Eli Lilly and Company, USA
SO Eur. Pat. Appl., 44 pp.
CODEN: EPXXDW
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 1609789	A1	20051228	EP 2004-380131	20040623
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, HR				
	WO 2006009741	A1	20060126	WO 2005-US21148	20050615
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	EP 1761520	A1	20070314	EP 2005-766569	20050615
	R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BA				

PRAI EP 2004-380131 A 20040623
US 2004-592539P P 20040730
EP 2004-380174 A 20040823
US 2004-622492P P 20041027
WO 2005-US21148 W 20050615
OS MARPAT 144:69825
RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 5 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2005:1311368 CAPLUS
DN 144:36261
TI Preparation of aroyl-O-piperidine derivatives as microsomal triglyceride transfer protein (MTP) and/or apoprotein B (ApoB) inhibitors useful in the treatment of dyslipidemia and related diseases
IN Guedat, Philippe; Collonges, Francois; Chevreuil, Olivier; Dumas, Herve; Denuault, Marie Noelle; Yvon, Stephane; Kane, Peter; Laiton, Julia; Robertson, Avril; Wendt, Bernd
PA Merck Sante, Fr.
SO Fr. Demande, 122 pp.
CODEN: FRXXBL
DT Patent
LA French
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	FR 2871463	A1	20051216	FR 2004-6345	20040611
	FR 2871463	B1	20060922		
	AU 2005251876	A1	20051222	AU 2005-251876	20050519
	CA 2569883	A1	20051222	CA 2005-2569883	20050519
	WO 2005121091	A1	20051222	WO 2005-EP5440	20050519

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

EP 1753721 A1 20070221 EP 2005-742232 20050519
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, LV

PRAI FR 2004-6345 A 20040611
WO 2005-EP5440 W 20050519

OS MARPAT 144:36261

RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 6 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN

AN 2005:1262710 CAPLUS

DN 144:22817

TI Preparation of phenyl or pyridinyl ureas as antagonists of P2Y1 receptors for the treatment of thromboembolic disorders

IN Chao, Hannguang J.; Tuerdi, Huji; Herpin, Timothy; Roberge, Jacques Yves; Liu, Yalei; Lawrence, R. Michael; Rehfuss, Robert P.; Clark, Charles G.; Qiao, Jennifer X.; Gungor, Timur; Lam, Patrick Y. S.; Wang, Tammy C.; Ruel, Rejean; L'Heureux, Alexandre L.; Thibeault, Carl; Bouthillier, Gilles; Schnur, Dora M.

PA Bristol-Myers Squibb Company, USA

SO PCT Int. Appl., 343 pp.
CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 2

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2005113511	A1	20051201	WO 2005-US16422	20050511
WO 2005113511	A9	20060202		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2005261244	A1	20051124	US 2005-126567	20050510
AU 2005245389	A1	20051201	AU 2005-245389	20050511
US 2005267119	A1	20051201	US 2005-126915	20050511
EP 1751113	A1	20070214	EP 2005-747470	20050511

R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, HR, LV, MK, YU

NO 2006005534	A	20061205	NO 2006-5534	20061130
PRAI US 2004-570288P	P	20040512		
US 2005-665735P	P	20050328		
US 2005-665817P	P	20050328		

	US 2005-126567	A	20050510
	WO 2005-US16422	W	20050511

OS MARPAT 144:22817
RE.CNT 15 THERE ARE 15 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 7 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2005:1260610 CAPLUS
DN 144:22946
TI Preparation of nitrogen-heteroaryl-containing protein kinase modulators for use against cancer and other diseases
IN Geuns-Meyer, Stephanie D.; Hodous, Brian L.; Chaffee, Stuart C.; Tempest, Paul A.; Olivieri, Philip R.; Johnson, Rebecca E.; Albrecht, Brian K.; Patel, Vinod F.; Cee, Victor J.; Kim, Joseph L.; Bellon, Steven; Zhu, Xiaotian; Cheng, Yuan; Xi, Ning; Romero, Karina; Nguyen, Hanh Nho; Deak, Holly L.
PA Amgen Inc., USA
SO PCT Int. Appl., 540 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2005113494	A2	20051201	WO 2005-US16346	20050509
	WO 2005113494	A3	20060316		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	AU 2005245386	A1	20051201	AU 2005-245386	20050509
	CA 2564355	A1	20051201	CA 2005-2564355	20050509
	US 2006009453	A1	20060112	US 2005-126000	20050509
	EP 1751136	A2	20070214	EP 2005-779977	20050509
	R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, LV, MK, YU				
PRAI	US 2004-569193P	P	20040507		
	WO 2005-US16346	W	20050509		

OS MARPAT 144:22946

L8 ANSWER 8 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2005:962206 CAPLUS
DN 143:266944
TI Preparation of heteroarylphenylurea derivatives as Raf inhibitors
IN Oikawa, Nobuhiro; Mizuguchi, Eisaku; Morikami, Kenji; Shimma, Nobuo; Ishii, Nobuya; Tsukaguchi, Toshiyuki; Ozawa, Sawako
PA Chugai Seiyaku Kabushiki Kaisha, Japan
SO PCT Int. Appl., 296 pp.
CODEN: PIXXD2
DT Patent
LA Japanese
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2005080330	A1	20050901	WO 2005-JP2923	20050223

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

EP 1724258 A1 20061122 EP 2005-719431 20050223
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR

PRAI JP 2004-47037 A 20040223
JP 2004-248856 A 20040827
WO 2005-JP2923 W 20050223

OS MARPAT 143:266944
RE.CNT 36 THERE ARE 36 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 9 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2004:1037107 CAPLUS
DN 142:23304
TI Preparation of pyrazoloquinazolines as inhibitors of protein kinases such as Aurora2 for the treatment of proliferative disorders such as cancer, Alzheimer's disease, and autoimmune diseases
IN Traquandi, Gabriella; Brasca, Maria Gabriella; D'Alessio, Roberto; Polucci, Paolo; Roletto, Fulvia; Vulpetti, Anna; Pevarello, Paolo; Panzeri, Achille; Quartieri, Francesca; Ferguson, Ron; Vianello, Paola; Fancelli, Daniele
PA Pharmacia Italia S.A., Italy
SO PCT Int. Appl., 226 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2004104007	A1	20041202	WO 2004-EP50612	20040427
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2004240772	A1	20041202	AU 2004-240772	20040427
CA 2526578	A1	20041202	CA 2004-2526578	20040427
EP 1636236	A1	20060322	EP 2004-741483	20040427
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK				
BR 2004010563	A	20060620	BR 2004-10563	20040427
CN 1826343	A	20060830	CN 2004-80021075	20040427
JP 2007502851	T	20070215	JP 2006-530168	20040427
NO 2005005496	A	20060214	NO 2005-5496	20051121
PRAI US 2003-472661P	P	20030522		
WO 2004-EP50612	W	20040427		

OS MARPAT 142:23304
RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 10 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2004:927173 CAPLUS
DN 141:395422
TI Preparation of N-[(piperidinyloxy)phenyl]-, N-[(piperidinyloxy)pyridinyl]-, N-[(piperidinylsulfanyl)phenyl]-, and N-[(piperidinylsulfanyl)pyridinyl] amides as 5-HT1F agonists for treatment of migraine
IN Blanco-Pillado, Maria-Jesus; Benesh, Dana Rae; Filla, Sandra Ann; Hudziak, Kevin John; Mathes, Brian Michael; Kohlman, Daniel Timothy; Ying, Bai-Ping; Zhang, Deyi; Xu, Yao-Chang
PA Eli Lilly and Company, USA
SO PCT Int. Appl., 186 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004094380	A1	20041104	WO 2004-US9283	20040414
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	AU 2004232799	A1	20041104	AU 2004-232799	20040414
	CA 2518839	A1	20041104	CA 2004-2518839	20040414
	EP 1626958	A1	20060222	EP 2004-759769	20040414
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK				
	BR 2004009211	A	20060328	BR 2004-9211	20040414
	CN 1777584	A	20060524	CN 2004-80010411	20040414
	JP 2006523692	T	20061019	JP 2006-509337	20040414
	US 2006211734	A1	20060921	US 2005-552131	20051011
PRAI	US 2003-464396P	P	20030418		
	WO 2004-US9283	A	20040414		

OS MARPAT 141:395422
RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 11 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2004:120834 CAPLUS
DN 140:181466
TI Preparation of resorcinol derivatives as peroxisome proliferator-activated receptor (PPAR) γ-agonists
IN Shibata, Tomoyuki; Wada, Kunio; Nakamura, Yuji; Araki, Kazushi
PA Sankyo Company, Limited, Japan
SO PCT Int. Appl., 261 pp.
CODEN: PIXXD2
DT Patent
LA Japanese
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004013109	A1	20040212	WO 2003-JP9834	20030801
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,				

LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

	AU 2003254795	A1	20040223	AU 2003-254795	20030801
	JP 2004123711	A	20040422	JP 2003-205222	20030801
PRAI	JP 2002-225980	A	20020802		
	WO 2003-JP9834	W	20030801		

OS MARPAT 140:181466

L8 ANSWER 12 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2003:950057 CAPLUS
DN 140:16647
TI Preparation of 2-aminopyridine-3-carboxamides as remedies for angiogenesis mediated diseases
IN Askew, Benny; Adams, Jeffrey; Booker, Shon; Chen, Guoqing; DiPietro, Lucian V.; Elbaum, Daniel; Germain, Julie; Geuns-Meyer, Stephanie D.; Habgood, Gregory J.; Handley, Michael; Huang, Qi; Kim, Tae-seong; Li, Aiwen; Nishimura, Nobuko; Nomak, Rana; Patel, Vinod F.; Riahi, Babak; Kim, Joseph L.; Xi, Ning; Yang, Kevin; Yuan, Chester Chenguang
PA Amgen Inc., USA
SO U.S. Pat. Appl. Publ., 252 pp., Cont.-in-part of U.S. Ser. No. 46,681.
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2003225106	A1	20031204	US 2002-197974	20020717
	US 6878714	B2	20050412		
	US 2003125339	A1	20030703	US 2002-46681	20020110
	US 6995162	B2	20060207		
	AT 361288	T	20070515	AT 2002-717325	20020111
	ZA 2003005197	A	20040319	ZA 2003-5197	20030704
	CA 2492100	A1	20040122	CA 2003-2492100	20030715
	WO 2004007458	A1	20040122	WO 2003-US22417	20030715
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	AU 2003252011	A1	20040202	AU 2003-252011	20030715
	EP 1537084	A1	20050608	EP 2003-764794	20030715
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	JP 2006501195	T	20060112	JP 2004-521959	20030715
	BG 108012	A	20041130	BG 2003-108012	20030721
	US 2005261313	A1	20051124	US 2004-14184	20041215
	US 2006040956	A1	20060223	US 2005-234713	20050923
	AU 2006200437	A1	20060223	AU 2006-200437	20060201
PRAI	US 2001-261339P	P	20010112		
	US 2001-323764P	P	20010919		
	US 2002-46681	A2	20020110		
	AU 2002-248340	A3	20020111		
	US 2002-197974	A	20020717		
	WO 2003-US22417	W	20030715		

OS MARPAT 140:16647
RE.CNT 39 THERE ARE 39 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 13 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2003:855655 CAPLUS
DN 139:350636
TI Preparation of amino heteroaryl amides for use in pharmaceutical compositions for the treatment of angiogenesis mediated diseases such as cancer
IN Patel, Vinod F.; Askew, Benny; Booker, Shon; Chen, Guoqing; Dipietro, Lucian V.; Germain, Julie; Habgood, Gregory J.; Huang, Qi; Kim, Tae-seong; Li, Aiwen; Nishimura, Nobuko; Nomak, Rana; Riahi, Babak; Yuan, Chester Chenguang; Elbaum, Daniel
PA Amgen Inc., USA
SO U.S. Pat. Appl. Publ., 148 pp., Cont.-in-part of U.S. Ser. No. 46,622.
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2003203922	A1	20031030	US 2002-197918	20020717
	US 7102009	B2	20060905		
	US 2003195230	A1	20031016	US 2002-46622	20020110
	US 7105682	B2	20060912		
	CN 1538836	A	20041020	CN 2002-806467	20020111
	ZA 2003005198	A	20040630	ZA 2003-5198	20030704
	CA 2492045	A1	20040122	CA 2003-2492045	20030715
	WO 2004007481	A2	20040122	WO 2003-US22275	20030715
	WO 2004007481	A3	20040219		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	AU 2003263784	A1	20040202	AU 2003-263784	20030715
	EP 1562933	A2	20050817	EP 2003-764755	20030715
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	JP 2006502118	T	20060119	JP 2004-521922	20030715
	US 2006194848	A1	20060831	US 2006-417329	20060502
PRAI	US 2001-261882P	P	20010112		
	US 2001-323808P	P	20010919		
	US 2002-46622	A2	20020110		
	US 2002-197918	A	20020717		
	WO 2003-US22275	W	20030715		

OS MARPAT 139:350636
RE.CNT 84 THERE ARE 84 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 14 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2002:965133 CAPLUS
DN 138:39277
TI Preparation of N-thiazolyl-N'-pyridyl ureas as antitumor agents
IN Askew, Benny C.; De Morin, Frenel F.; Hague, Andrew; Laber, Ellen; Li, Aiwen; Liu, Gang; Lopez, Patricia; Nomak, Rana; Santora, Vincent; Tegley, Christopher; Yang, Kevin
PA Amgen, Inc., USA

SO U.S. Pat. Appl. Publ., 129 pp., Cont.-in-part of U. S. Ser. No. 930,753.
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 2

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 2002193405	A1	20021219	US 2002-77124	20020215
US 6645990	B2	20031111		
US 2002173507	A1	20021121	US 2001-930753	20010814
EP 1619184	A2	20060125	EP 2005-15480	20010815
EP 1619184	A3	20060201		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
AT 320426	T	20060415	AT 2001-964009	20010815
ES 2260277	T3	20061101	ES 2001-1964009	20010815
CA 2476411	A1	20030828	CA 2003-2476411	20030213
WO 2003070727	A1	20030828	WO 2003-US4537	20030213
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2003215231	A1	20030909	AU 2003-215231	20030213
EP 1483263	A1	20041208	EP 2003-711046	20030213
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
JP 2006509715	T	20060323	JP 2003-569634	20030213
US 2004039029	A1	20040226	US 2003-631423	20030730
US 7196104	B2	20070327		
US 2004044044	A1	20040304	US 2003-632044	20030730
PRAI US 2000-225793P	P	20000815		
US 2001-930753	A2	20010814		
EP 2001-964009	A3	20010815		
US 2002-77124	A	20020215		
WO 2003-US4537	W	20030213		

OS MARPAT 138:39277

L8 ANSWER 15 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2002:736252 CAPLUS
DN 137:263031
TI Preparation of 5-substituted imidazolidine-2,4-diones as metalloproteinase inhibitors
IN Eriksson, Anders; Lepistoe, Matti; Lundkvist, Michael; Munck Af Rosenschoeld, Magnus; Zlatoidsky, Pavol
PA Astrazeneca AB, Swed.
SO PCT Int. Appl., 153 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 6

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2002074767	A1	20020926	WO 2002-SE472	20020313
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,				

UA, UG, US, UZ, VN, YU, ZA, ZM, ZW
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

CA 2440630	A1	20020926	CA 2002-2440630	20020313
AU 2002237626	A1	20021003	AU 2002-237626	20020313
AU 2002237626	B2	20070517		
EE 200300445	A	20031215	EE 2003-445	20020313
EP 1370556	A1	20031217	EP 2002-704031	20020313
EP 1370556	B1	20060719		

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR

BR 2002008104	A	20040302	BR 2002-8104	20020313
CN 1509272	A	20040630	CN 2002-809788	20020313
CN 1509286	A	20040630	CN 2002-809915	20020313
CN 1509276	A	20040630	CN 2002-810093	20020313
JP 2004527515	T	20040909	JP 2002-573776	20020313
HU 200400327	A2	20050128	HU 2004-327	20020313
NZ 528106	A	20050324	NZ 2002-528106	20020313
EP 1676846	A2	20060705	EP 2006-8158	20020313
EP 1676846	A3	20060726		

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR

AT 333454	T	20060815	AT 2002-704031	20020313
RU 2288228	C2	20061127	RU 2003-127734	20020313
CN 1962641	A	20070516	CN 2006-10106152	20020313
IN 2003MN00805	A	20050318	IN 2003-MN805	20030827
ZA 2003006731	A	20041129	ZA 2003-6731	20030828
ZA 2003006732	A	20041129	ZA 2003-6732	20030828
ZA 2003006734	A	20041129	ZA 2003-6734	20030828
ZA 2003006737	A	20041129	ZA 2003-6737	20030828
NO 2003004045	A	20031110	NO 2003-4045	20030912
US 2004127528	A1	20040701	US 2004-471900	20040114
HK 1059932	A1	20061222	HK 2004-102796	20040421
PRAI SE 2001-902	A	20010315		
CN 2002-810093	A3	20020313		
EP 2002-704031	A3	20020313		
WO 2002-SE472	W	20020313		

OS MARPAT 137:263031
RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 16 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2002:736236 CAPLUS
DN 137:247696
TI Preparation of 5-substituted imidazolidine-2,4-diones as metalloproteinase inhibitors
IN Eriksson, Anders; Lepistoe, Matti; Lundkvist, Michael; Munck Af Rosenschoeld, Magnus; Zlatoidsky, Pavol
PA Astrazeneca AB, Swed.
SO PCT Int. Appl., 300 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 6

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2002074750	A1	20020926	WO 2002-SE475	20020313

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,

UA, UG, US, UZ, VN, YU, ZA, ZM, ZW
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

	CA 2440632	A1	20020926	CA 2002-2440632	20020313
	AU 2002237629	A1	20021003	AU 2002-237629	20020313
	EE 200300439	A	20031215	EE 2003-439	20020313
	EP 1370536	A1	20031217	EP 2002-704034	20020313
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	BR 2002008105	A	20040309	BR 2002-8105	20020313
	CN 1509275	A	20040630	CN 2002-810041	20020313
	HU 200400206	A2	20040830	HU 2004-206	20020313
	JP 2004527511	T	20040909	JP 2002-573759	20020313
	EP 1676846	A2	20060705	EP 2006-8158	20020313
	EP 1676846	A3	20060726		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	CN 1962641	A	20070516	CN 2006-10106152	20020313
	IN 2003MN00800	A	20050318	IN 2003-MN800	20030827
	NO 2003004025	A	20031113	NO 2003-4025	20030911
	US 2004147573	A1	20040729	US 2003-471808	20030912
PRAI	SE 2001-902	A	20010315		
	SE 2001-903	A	20010315		
	CN 2002-810093	A3	20020313		
	EP 2002-704031	A3	20020313		
	WO 2002-SE475	W	20020313		

OS MARPAT 137:247696
RE.CNT 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 17 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2002:676007 CAPLUS
DN 137:216945
TI Preparation of substituted 2-(1H-indazol-6-ylamino)nicotinamides for treating KDR-related diseases
IN Chen, Guoqing; Adams, Jeffrey; Bemis, Jean; Croghan, Michael; Dipietro, Lucian; Dominguez, Celia; Elbaum, Daniel; Germain, Julie; Huang, Qi; Kim, Joseph L.; Ouyang, Xiaohu; Patel, Vinod F.; Smith, Leon M.; Tasker, Andrew; Xi, Ning; Xu, Shimin; Yuan, Chester Chenguang; Kim, Tae-Seong
PA Amgen Inc., USA
SO PCT Int. Appl., 395 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002068406	A2	20020906	WO 2002-US3064	20020111
	WO 2002068406	A3	20030424		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	US 2003195230	A1	20031016	US 2002-46622	20020110
	US 7105682	B2	20060912		
	CA 2434178	A1	20020906	CA 2002-2434178	20020111
	AU 2002253890	A1	20020912	AU 2002-253890	20020111

	HU 200302719	A2	20031128	HU 2003-2719	20020111
	EE 200300325	A	20031215	EE 2003-325	20020111
	JP 2004527499	T	20040909	JP 2002-567920	20020111
	CN 1538836	A	20041020	CN 2002-806467	20020111
	EP 1467721	A2	20041020	EP 2002-723086	20020111
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	ZA 2003005198	A	20040630	ZA 2003-5198	20030704
	BG 108013	A	20040430	BG 2003-108013	20030721
	US 2006194848	A1	20060831	US 2006-417329	20060502
PRAI	US 2001-261882P	P	20010112		
	US 2001-323808P	P	20010919		
	US 2002-46622	A	20020110		
	WO 2002-US3064	W	20020111		

OS MARPAT 137:216945

L8 ANSWER 18 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2002:658116 CAPLUS
DN 137:201332
TI Preparation of heterocyclylalkylamine derivatives as remedies for angiogenesis mediated diseases
IN Chen, Guoqing; Adams, Jeffrey; Bemis, Jean; Booker, Shon; Cai, Guolin; Croghan, Michael; DiPietro, Lucian; Dominguez, Celia; Elbaum, Daniel; Germain, Julie; Geuns-Meyer, Stephanie; Handley, Michael; Huang, Qi; Kim, Joseph L.; Kim, Tae-seong; Kiselyov, Alexander; Ouyang, Xiaohu; Patel, Vinod F.; Smith, Leon M.; Stec, Markian; Tasker, Andrew; Xi, Ning; Xu, Shimin; Yuan, Chester Chenguang
PA Amgen Inc., USA
SO PCT Int. Appl., 502 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002066470	A1	20020829	WO 2002-US743	20020111
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	US 2003125339	A1	20030703	US 2002-46681	20020110
	US 6995162	B2	20060207		
	CA 2434277	A1	20020829	CA 2002-2434277	20020111
	AU 2002248340	A1	20020904	AU 2002-248340	20020111
	BR 2002006435	A	20030923	BR 2002-6435	20020111
	EP 1358184	A1	20031105	EP 2002-717325	20020111
	EP 1358184	B1	20070502		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	HU 200302598	A2	20031128	HU 2003-2598	20020111
	EE 200300324	A	20031215	EE 2003-324	20020111
	JP 2004531484	T	20041014	JP 2002-565984	20020111
	NZ 526868	A	20050429	NZ 2002-526868	20020111
	CN 1671700	A	20050921	CN 2002-806202	20020111
	AT 361288	T	20070515	AT 2002-717325	20020111
	ZA 2003005197	A	20040319	ZA 2003-5197	20030704
	NO 2003003181	A	20030911	NO 2003-3181	20030711
	IN 2003CN01070	A	20050422	IN 2003-CN1070	20030711
	BG 108012	A	20041130	BG 2003-108012	20030721

	US 2006040956	A1	20060223	US 2005-234713	20050923
	AU 2006200437	A1	20060223	AU 2006-200437	20060201
PRAI	US 2001-261339P	P	20010112		
	US 2001-323764P	P	20010919		
	US 2002-46681	A	20020110		
	AU 2002-248340	A3	20020111		
	WO 2002-US743	W	20020111		

OS MARPAT 137:201332
RE.CNT 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 19 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2001:472725 CAPLUS
DN 135:76897
TI Synthesis and use of substituted piperidine and piperazine derivatives (e.g. N-(sulfonyl)aryl, N-alkylcarboxamido piperazines) as antagonists of the P2X7 receptor
IN Meghani, Premji; Bennion, Colin
PA Astrazeneca AB, Swed.
SO PCT Int. Appl., 156 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001046200	A1	20010628	WO 2000-SE2580	20001218
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	CA 2394095	A1	20010628	CA 2000-2394095	20001218
	BR 2000016543	A	20020917	BR 2000-16543	20001218
	EP 1242427	A1	20020925	EP 2000-989102	20001218
	EP 1242427	B1	20030813		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	JP 2003518126	T	20030603	JP 2001-547110	20001218
	AT 247123	T	20030815	AT 2000-989102	20001218
	NZ 519498	A	20040227	NZ 2000-519498	20001218
	AU 776592	B2	20040916	AU 2001-25648	20001218
	ZA 2002004307	A	20030829	ZA 2002-4307	20020529
	US 2003013721	A1	20030116	US 2002-168094	20020617
	US 6969713	B2	20051129		
	NO 2002003037	A	20020822	NO 2002-3037	20020621
	US 2005272745	A1	20051208	US 2005-125335	20050510
PRAI	SE 1999-4738	A	19991222		
	WO 2000-SE2580	W	20001218		
	US 2002-168094	A1	20020617		

OS MARPAT 135:76897
RE.CNT 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 20 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2000:842122 CAPLUS
DN 134:17318
TI Preparation of substituted 2-phenylamino-N-phenylacetamides with immunosuppressing activity
IN Furber, Mark; Luker, Timothy Jon; Mortimore, Michael Paul; Thorne, Philip;

Meghani, Premji
PA Astrazeneca AB, Swed.
SO PCT Int. Appl., 68 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2000071529	A1	20001130	WO 2000-GB1943	20000522
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
CA 2372580	A1	20001130	CA 2000-2372580	20000522
BR 2000010716	A	20020213	BR 2000-10716	20000522
EP 1185522	A1	20020313	EP 2000-931406	20000522
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
JP 2003500399	T	20030107	JP 2000-619786	20000522
NZ 515282	A	20040130	NZ 2000-515282	20000522
AU 778305	B2	20041125	AU 2000-49362	20000522
US 6555541	B1	20030429	US 2000-583000	20000710
ZA 2001009091	A	20030203	ZA 2001-9091	20011102
NO 2001005665	A	20020124	NO 2001-5665	20011120
NO 322166	B1	20060821		
PRAI SE 1999-1875	A	19990525		
WO 2000-GB1943	W	20000522		

OS MARPAT 134:17318
RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 21 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2000:742083 CAPLUS
DN 133:309908
TI Preparation of piperazinyladamantylmethylbenzamides and related compounds as P2X7 receptor antagonists.
IN Alcaraz, Lilian; Furber, Mark; Mortimore, Michael
PA AstraZeneca AB, Swed.
SO PCT Int. Appl., 166 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2000061569	A1	20001019	WO 2000-SE663	20000406
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW				
RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
CA 2368829	A1	20001019	CA 2000-2368829	20000406
BR 2000009651	A	20020108	BR 2000-9651	20000406
EP 1171432	A1	20020116	EP 2000-919245	20000406
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				

	TR 200102911	T2	20020121	TR 2001-2911	20000406
	HU 200202214	A2	20021028	HU 2002-2214	20000406
	JP 2002541249	T	20021203	JP 2000-610843	20000406
	EE 200100525	A	20021216	EE 2001-525	20000406
	EE 4565	B1	20051215		
	NZ 514477	A	20030429	NZ 2000-514477	20000406
	AU 774526	B2	20040701	AU 2000-39947	20000406
	RU 2254333	C2	20050620	RU 2001-130140	20000406
	US 6492355	B1	20021210	US 2000-555489	20000601
	IN 2001MN01201	A	20050318	IN 2001-MN1201	20011001
	NO 2001004894	A	20011210	NO 2001-4894	20011008
	NO 321405	B1	20060508		
	ZA 2001008265	A	20030108	ZA 2001-8265	20011008
PRAI	SE 1999-1270	A	19990409		
	GB 2000-2330	A	20000201		
	WO 2000-SE663	W	20000406		

OS MARPAT 133:309908
RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 22 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 1998:745020 CAPLUS
DN 130:13850
TI Preparation of arylacetamide and arylurea derivatives as 5-HT1A, 5-HT1B, and 5-HT1D receptor antagonists.
IN Gaster, Laramie Mary; Wyman, Paul Adrian
PA Smithkline Beecham PLC, UK
SO PCT Int. Appl., 73 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9850346	A2	19981112	WO 1998-EP2263	19980414
	WO 9850346	A3	19990311		
	W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
	AU 9875267	A	19981127	AU 1998-75267	19980414
	ZA 9803243	A	19991018	ZA 1998-3243	19980417
PRAI	GB 1997-7874	A	19970418		
	GB 1998-1632	A	19980126		
	WO 1998-EP2263	W	19980414		

OS MARPAT 130:13850

L8 ANSWER 23 OF 23 CAPLUS COPYRIGHT 2007 ACS on STN
AN 1995:501323 CAPLUS
DN 122:265361
TI Preparation of 3-aryl-5-[(4-aryloxy- and -thiopiperidino)alkyl]oxazolidin-2-ones as nervous system agents
IN Pruecher, Helmut; Gottschlich, Rudolf; Bartoszyk, Gerd; Seyfried, Christoph
PA Merck Patent G.m.b.H., Germany
SO Eur. Pat. Appl., 18 pp.
CODEN: EPXXDW
DT Patent
LA German
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 635505	A1	19950125	EP 1994-110781	19940712
	EP 635505	B1	19971015		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE				
	DE 4324393	A1	19950126	DE 1993-4324393	19930721
	AT 159252	T	19971115	AT 1994-110781	19940712
	ES 2110660	T3	19980216	ES 1994-110781	19940712
	SK 281630	B6	20010611	SK 1994-852	19940714
	AU 9467536	A	19950202	AU 1994-67536	19940715
	AU 683886	B2	19971127		
	TW 401417	B	20000811	TW 1994-83106530	19940718
	CA 2128380	A1	19950122	CA 1994-2128380	19940719
	CA 2128380	C	20050412		
	CZ 284544	B6	19981216	CZ 1994-1738	19940719
	PL 177692	B1	20000131	PL 1994-304349	19940719
	NO 9402715	A	19950123	NO 1994-2715	19940720
	ZA 9405340	A	19950301	ZA 1994-5340	19940720
	JP 07070117	A	19950314	JP 1994-168105	19940720
	CN 1106008	A	19950802	CN 1994-107977	19940720
	CN 1055690	B	20000823		
	RU 2135495	C1	19990827	RU 1994-26079	19940720
	HU 71110	A2	19951128	HU 1994-2154	19940721
	HU 218912	B	20001228		
	US 5561145	A	19961001	US 1994-278210	19940721
PRAI	DE 1993-4324393	A	19930721		
OS	MARPAT 122:265361				

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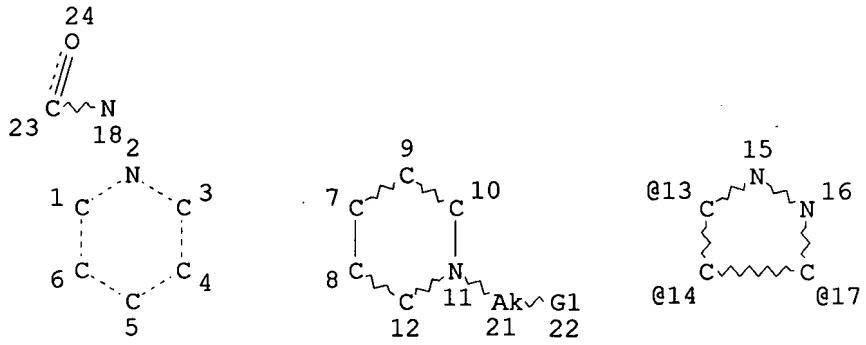

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DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
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NUMBER OF NODES IS 22

STEREO ATTRIBUTES: NONE

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SEARCH TIME: 00.00.01

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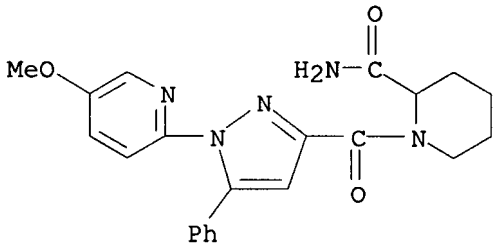
L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 2-Piperidinecarboxamide, 1-[[1-(5-methoxy-2-pyridinyl)-5-phenyl-1H-pyrazol-3-yl]carbonyl]- (9CI)
MF C22 H23 N5 O3



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

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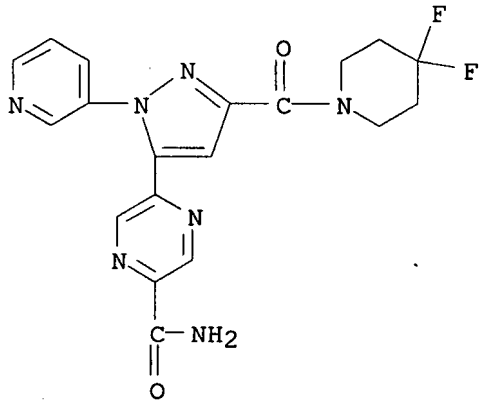
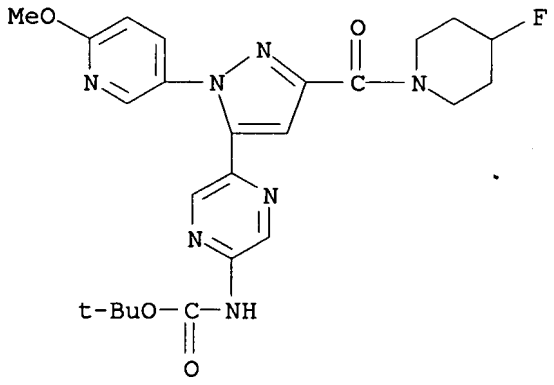
L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Pyrazinecarboxamide, 5-[3-[(4,4-difluoro-1-piperidinyl)carbonyl]-1-(3-pyridinyl)-1H-pyrazol-5-yl]- (9CI)
MF C19 H17 F2 N7 O2



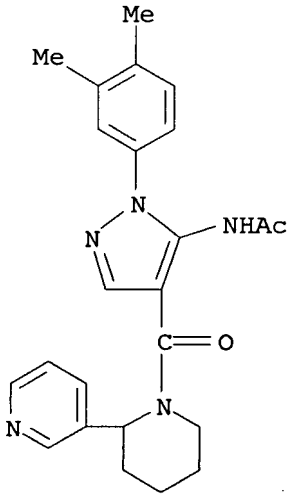
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Carbamic acid, [5-[3-[(4-fluoro-1-piperidinyl)carbonyl]-1-(6-methoxy-3-pyridinyl)-1H-pyrazol-5-yl]pyrazinyl]-, 1,1-dimethylethyl ester (9CI)
MF C24 H28 F N7 O4



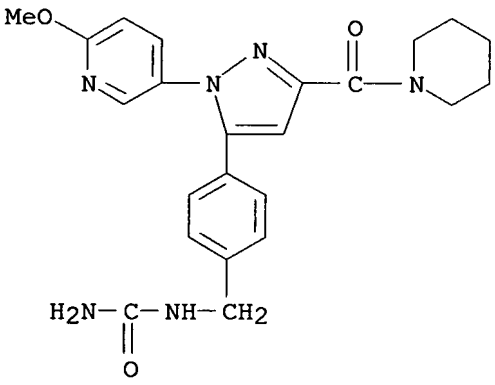
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Acetamide, N-[1-(3,4-dimethylphenyl)-4-[[2-(3-pyridinyl)-1-piperidinyl]carbonyl]-1H-pyrazol-5-yl]-
MF C24 H27 N5 O2



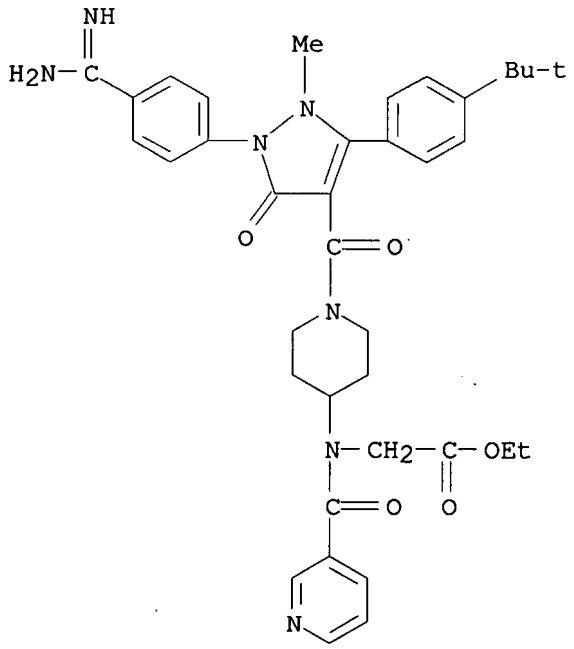
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Piperidine, 1-[[5-[4-[[(aminocarbonyl)amino]methyl]phenyl]-1-(6-methoxy-3-pyridinyl)-1H-pyrazol-3-yl]carbonyl]- (9CI)
MF C23 H26 N6 O3



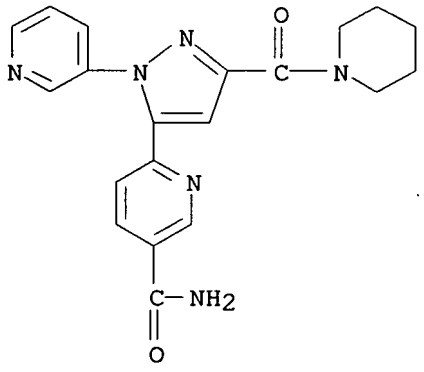
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Glycine, N-[1-[[2-[4-(aminoiminomethyl)phenyl]-5-[4-(1,1-dimethylethyl)phenyl]-2,3-dihydro-1-methyl-3-oxo-1H-pyrazol-4-yl]carbonyl]-4-piperidinyl]-N-(3-pyridinylcarbonyl)-, ethyl ester (9CI)
MF C37 H43 N7 O5
CI COM



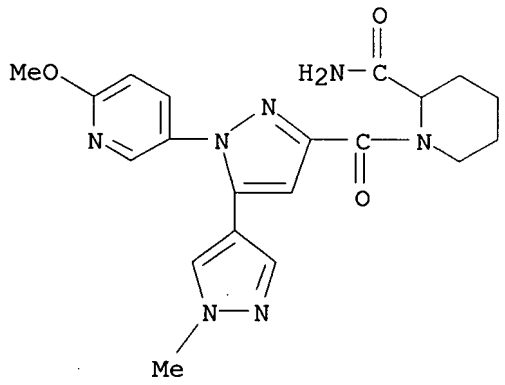
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 3-Pyridinecarboxamide, 6-[3-(1-piperidinylcarbonyl)-1-(3-pyridinyl)-1H-pyrazol-5-yl]- (9CI)
MF C20 H20 N6 O2



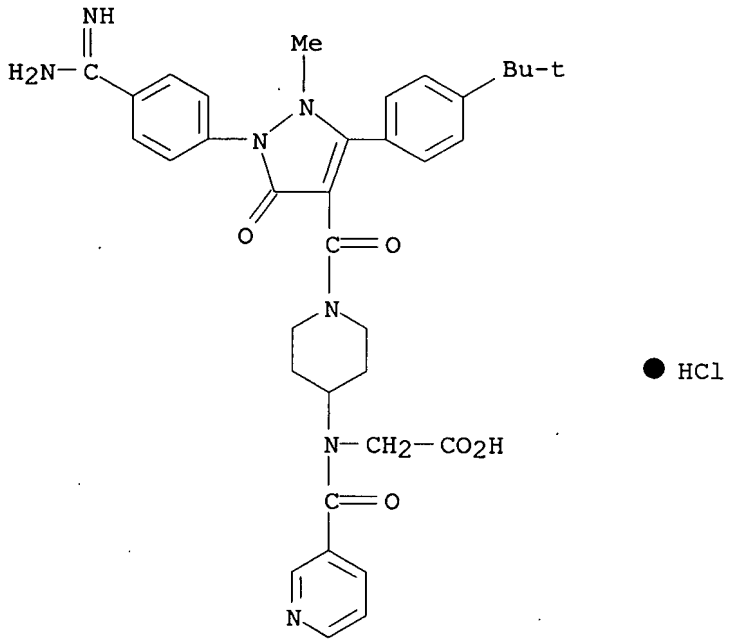
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 2-Piperidinecarboxamide, 1-[[1'-(6-methoxy-3-pyridinyl)-1-methyl[4,5'-bi-1H-pyrazol]-3'-yl]carbonyl]- (9CI)
MF C20 H23 N7 O3

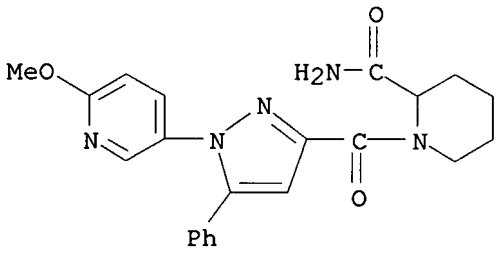


PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Glycine, N-[1-[[2-[4-(aminoiminomethyl)phenyl]-5-[4-(1,1-dimethylethyl)phenyl]-2,3-dihydro-1-methyl-3-oxo-1H-pyrazol-4-yl]carbonyl]-4-piperidinyl]-N-(3-pyridinylcarbonyl)-, monohydrochloride (9CI)
MF C35 H39 N7 O5 . Cl H



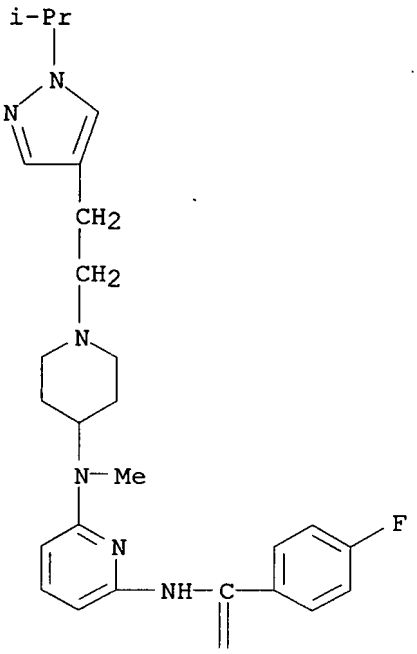
L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 2-Piperidinecarboxamide, 1-[[1-(6-methoxy-3-pyridinyl)-5-phenyl-1H-pyrazol-3-yl]carbonyl]- (9CI)
MF C22 H23 N5 O3



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Benzamide, 4-fluoro-N-[6-[methyl[1-[2-[1-(1-methylethyl)-1H-pyrazol-4-yl]ethyl]-4-piperidinyl]amino]-2-pyridinyl]-, monohydrochloride (9CI)
MF C26 H33 F N6 O . Cl H

PAGE 1-A



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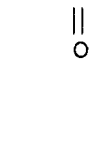
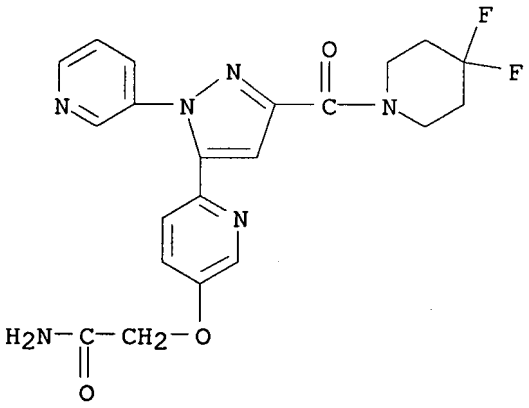


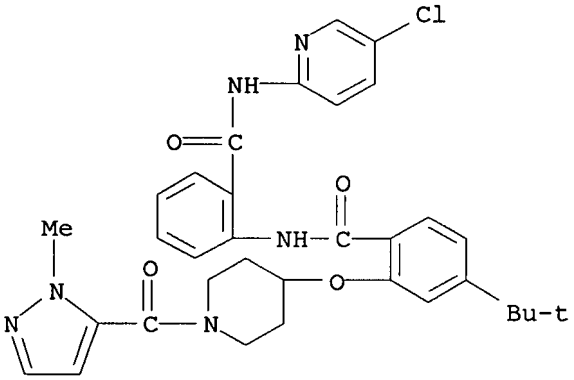
L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Acetamide, 2-[[6-[3-[(4,4-difluoro-1-piperidinyl)carbonyl]-1-(3-pyridinyl)-

1H-pyrazol-5-yl]-3-pyridinyl]oxy]- (9CI)
MF C21 H20 F2 N6 O3



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

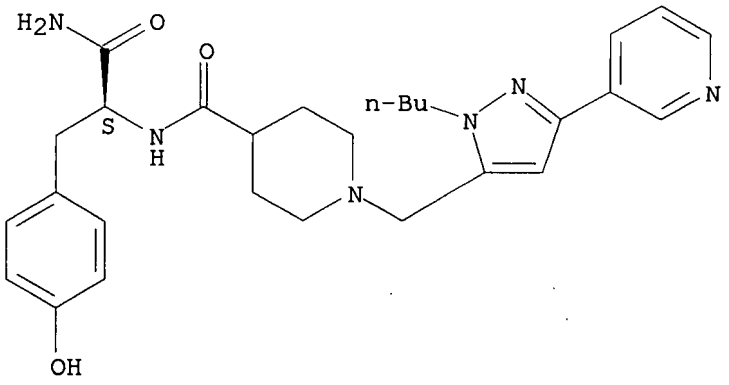
L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Benzamide, N-[2-[[(5-chloro-2-pyridinyl)amino]carbonyl]phenyl]-4-(1,1-dimethylethyl)-2-[[1-[(1-methyl-1H-pyrazol-5-yl)carbonyl]-4-piperidinyl]oxy]- (9CI)
MF C33 H35 Cl N6 O4



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

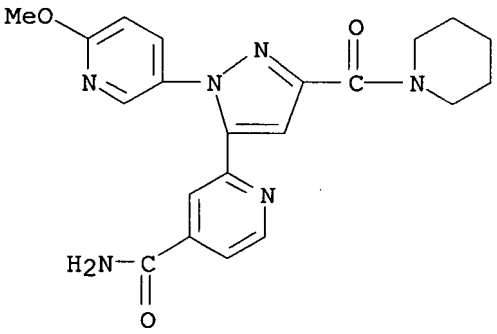
L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 4-Piperidinecarboxamide, N-[(1S)-2-amino-1-[(4-hydroxyphenyl)methyl]-2-oxoethyl]-1-[[1-butyl-3-(3-pyridinyl)-1H-pyrazol-5-yl]methyl]- (9CI)
MF C28 H36 N6 O3

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

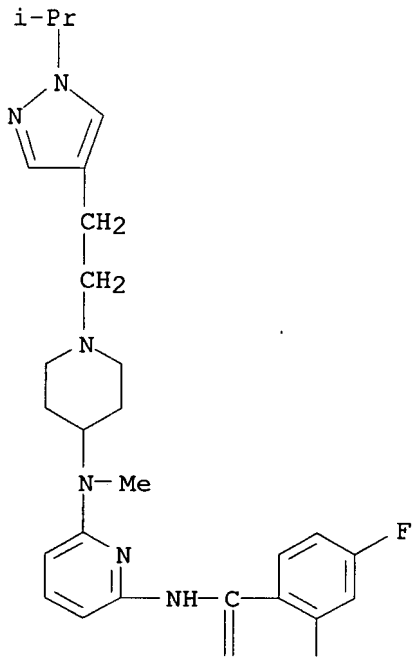
L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 4-Pyridinecarboxamide, 2-[1-(6-methoxy-3-pyridinyl)-3-(1-piperidinylcarbonyl)-1H-pyrazol-5-yl]- (9CI)
MF C21 H22 N6 O3



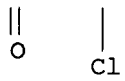
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Benzamide, 2-chloro-4-fluoro-N-[6-[methyl[1-[2-[1-(1-methylethyl)-1H-pyrazol-4-yl]ethyl]-4-piperidinyl]amino]-2-pyridinyl]- (9CI)
MF C26 H32 Cl F N6 O
CI COM

PAGE 1-A

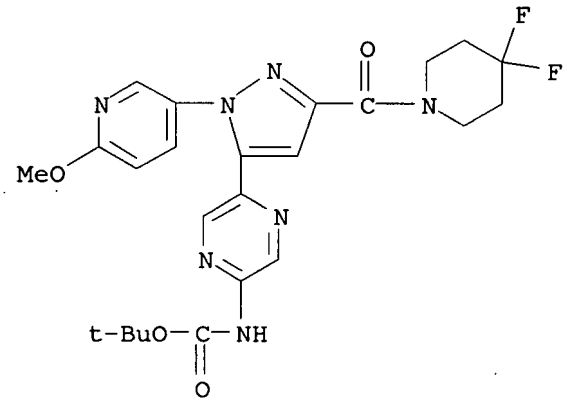


PAGE 2-A



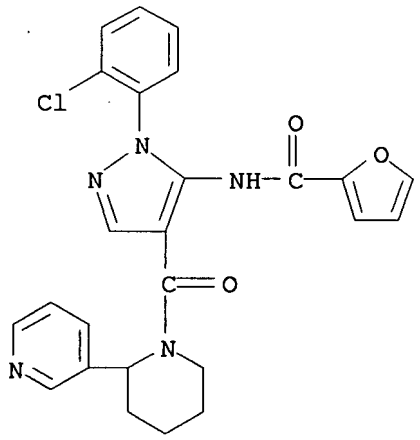
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Carbamic acid, [5-[3-[(4,4-difluoro-1-piperidinyl)carbonyl]-1-(6-methoxy-3-pyridinyl)-1H-pyrazol-5-yl]pyrazinyl]-, 1,1-dimethylethyl ester (9CI)
MF C24 H27 F2 N7 O4



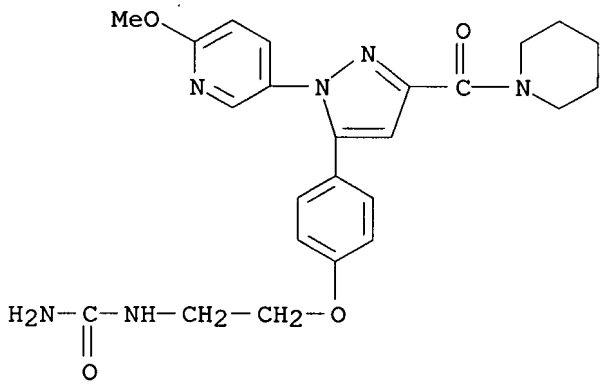
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 2-Furancarboxamide, N-[1-(2-chlorophenyl)-4-[[2-(3-pyridinyl)-1-piperidinyl]carbonyl]-1H-pyrazol-5-yl]-
MF C25 H22 Cl N5 O3



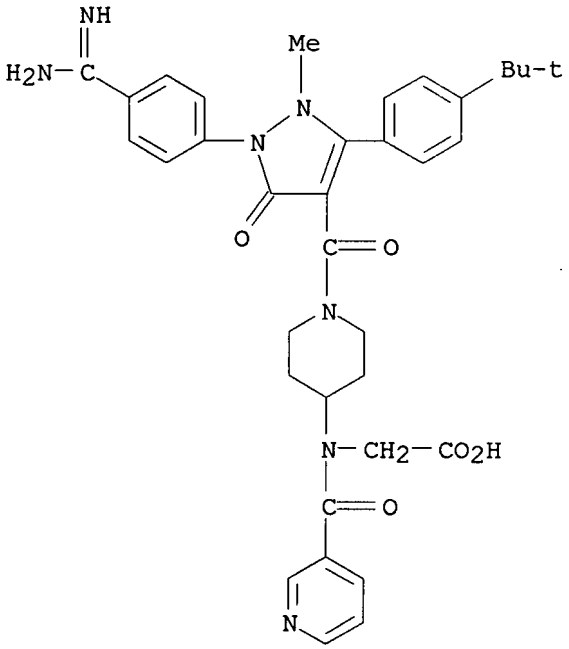
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Piperidine, 1-[[5-[4-[2-[(aminocarbonyl)amino]ethoxy]phenyl]-1-(6-methoxy-3-pyridinyl)-1H-pyrazol-3-yl]carbonyl]- (9CI)
MF C24 H28 N6 O4



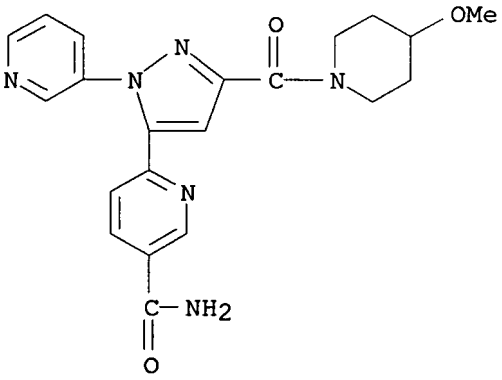
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Glycine, N-[1-[[2-[4-(aminoiminomethyl)phenyl]-5-[4-(1,1-dimethylethyl)phenyl]-2,3-dihydro-1-methyl-3-oxo-1H-pyrazol-4-yl]carbonyl]-4-piperidinyl]-N-(3-pyridinylcarbonyl)- (9CI)
MF C35 H39 N7 O5
CI COM



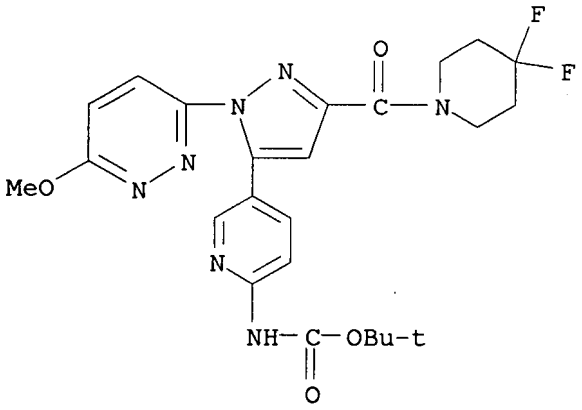
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 3-Pyridinecarboxamide, 6-[3-[(4-methoxy-1-piperidinyl)carbonyl]-1-(3-pyridinyl)-1H-pyrazol-5-yl]- (9CI)
MF C21 H22 N6 O3



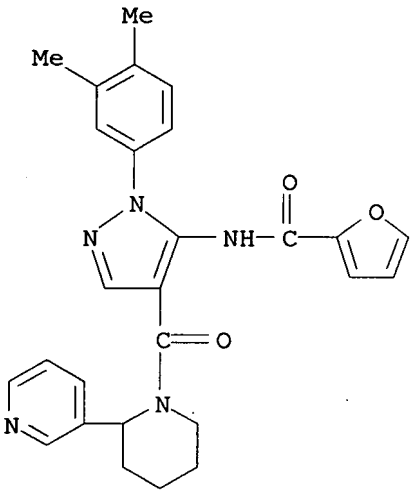
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Carbamic acid, [5-[3-[(4,4-difluoro-1-piperidinyl)carbonyl]-1-(6-methoxy-3-pyridazinyl)-1H-pyrazol-5-yl]-2-pyridinyl]-, 1,1-dimethylethyl ester (9CI)
MF C24 H27 F2 N7 O4



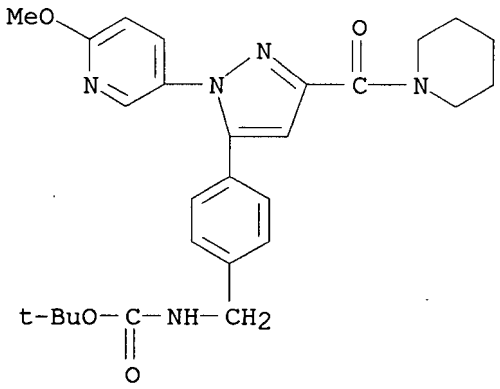
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 2-Furancarboxamide, N-[1-(3,4-dimethylphenyl)-4-[[2-(3-pyridinyl)-1-piperidinyl]carbonyl]-1H-pyrazol-5-yl]-
MF C27 H27 N5 O3



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Carbamic acid, [[4-[1-(6-methoxy-3-pyridinyl)-3-(1-piperidinylcarbonyl)-1H-pyrazol-5-yl]phenyl]methyl]-, 1,1-dimethylethyl ester (9CI)
MF C27 H33 N5 O4



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Benzamide, 2,4,6-trifluoro-N-[6-[[1-[2-[1-(1-methylethyl)-1H-pyrazol-4-yl]ethyl]-4-piperidinyl]oxy]-2-pyridinyl]-, monohydrochloride (9CI)
MF C25 H28 F3 N5 O2 . Cl H

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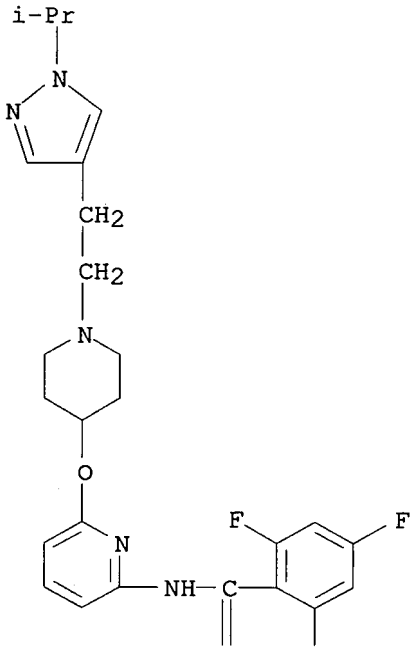

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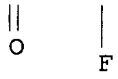
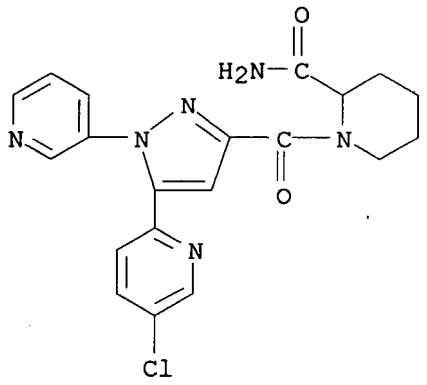


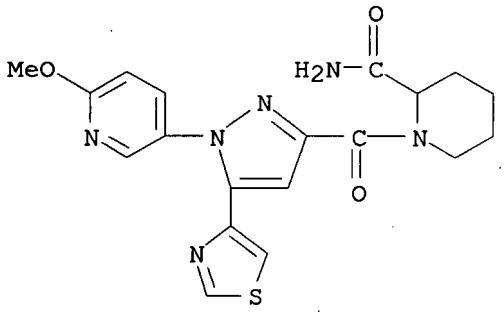
● HCl

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 2-Piperidinecarboxamide, 1-[[5-(5-chloro-2-pyridinyl)-1-(3-pyridinyl)-1H-pyrazol-3-yl]carbonyl]- (9CI)
MF C20 H19 Cl N6 O2



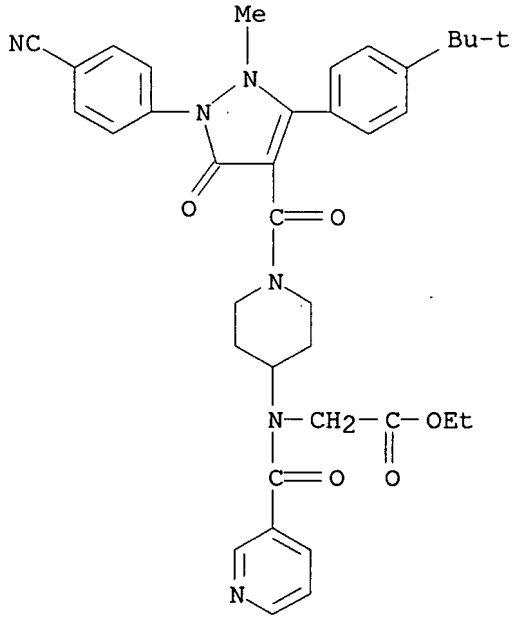
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 2-Piperidinecarboxamide, 1-[[1-(6-methoxy-3-pyridinyl)-5-(4-thiazolyl)-1H-pyrazol-3-yl]carbonyl]- (9CI)
MF C19 H20 N6 O3 S



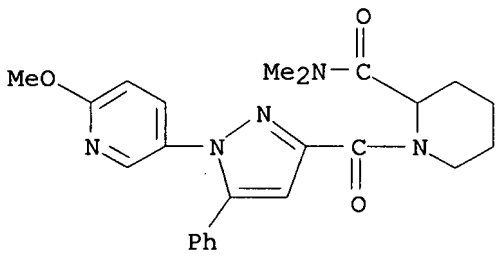
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Glycine, N-[1-[[2-(4-cyanophenyl)-5-[4-(1,1-dimethylethyl)phenyl]-2,3-dihydro-1-methyl-3-oxo-1H-pyrazol-4-yl]carbonyl]-4-piperidinyl]-N-(3-pyridinylcarbonyl)-, ethyl ester (9CI)
MF C37 H40 N6 O5



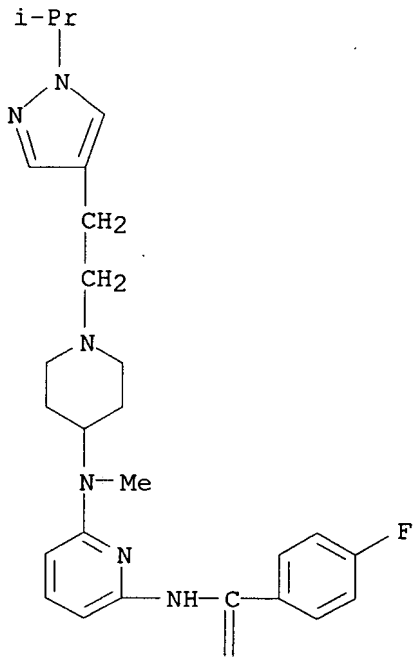
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 2-Piperidinecarboxamide, 1-[[1-(6-methoxy-3-pyridinyl)-5-phenyl-1H-pyrazol-3-yl]carbonyl]-N,N-dimethyl- (9CI)
MF C24 H27 N5 O3



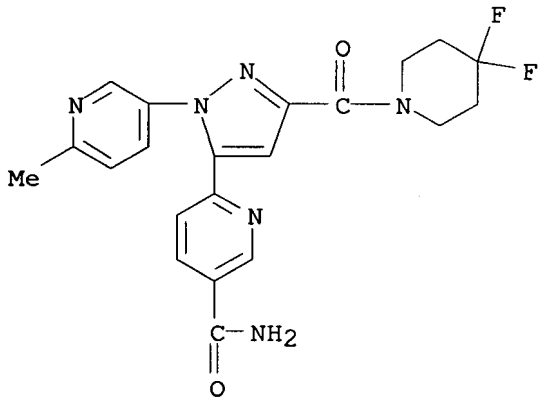
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Benzamide, 4-fluoro-N-[6-[methyl[1-[2-[1-(1-methylethyl)-1H-pyrazol-4-yl]ethyl]-4-piperidinyl]amino]-2-pyridinyl]- (9CI)
MF C26 H33 F N6 O
CI COM



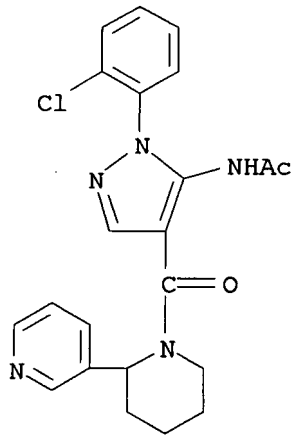
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 3-Pyridinecarboxamide, 6-[3-[(4,4-difluoro-1-piperidinyl)carbonyl]-1-(6-methyl-3-pyridinyl)-1H-pyrazol-5-yl]- (9CI)
MF C21 H20 F2 N6 O2



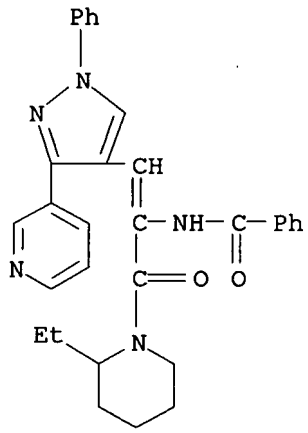
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Acetamide, N-[1-(2-chlorophenyl)-4-[[2-(3-pyridinyl)-1-piperidinyl]carbonyl]-1H-pyrazol-5-yl]-
MF C22 H22 Cl N5 O2



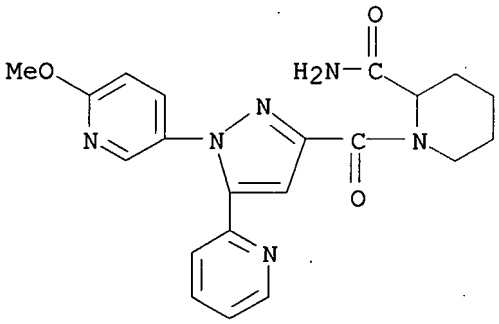
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Benzamide, N-[1-[(2-ethyl-1-piperidinyl)carbonyl]-2-[1-phenyl-3-(3-pyridinyl)-1H-pyrazol-4-yl]ethenyl]- (9CI)
MF C31 H31 N5 O2



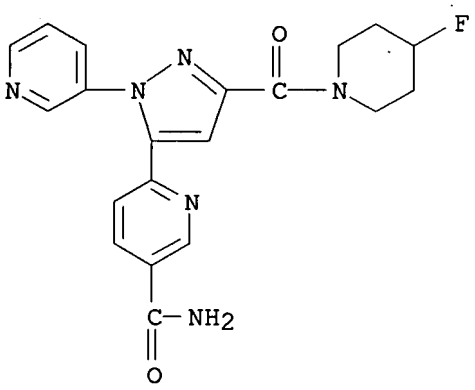
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 2-Piperidinecarboxamide, 1-[[1-(6-methoxy-3-pyridinyl)-5-(2-pyridinyl)-1H-pyrazol-3-yl]carbonyl]- (9CI)
MF C21 H22 N6 O3



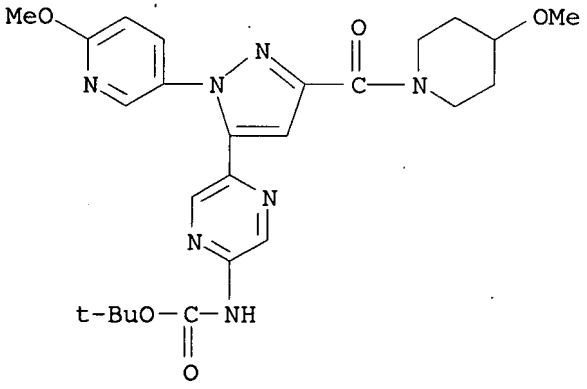
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 3-Pyridinecarboxamide, 6-[3-[(4-fluoro-1-piperidinyl)carbonyl]-1-(3-pyridinyl)-1H-pyrazol-5-yl]- (9CI)
MF C20 H19 F N6 O2



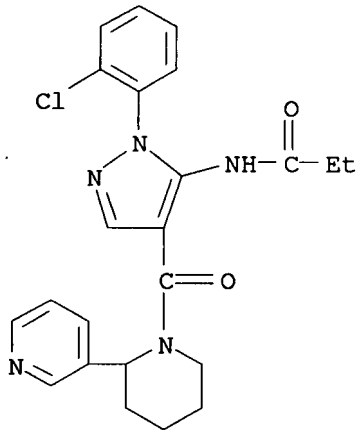
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Carbamic acid, [5-[3-[(4-methoxy-1-piperidinyl)carbonyl]-1-(6-methoxy-3-pyridinyl)-1H-pyrazol-5-yl]pyrazinyl]-, 1,1-dimethylethyl ester (9CI)
MF C25 H31 N7 O5



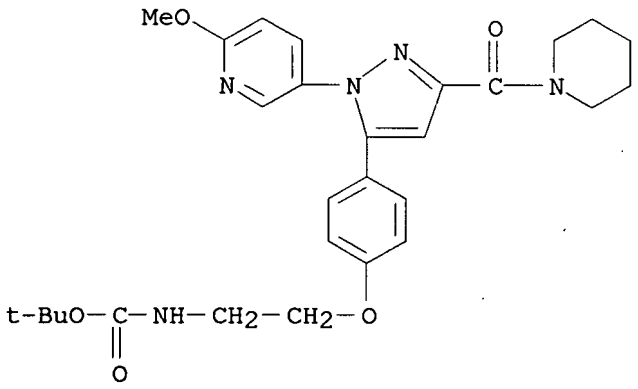
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Propanamide, N-[1-(2-chlorophenyl)-4-[[2-(3-pyridinyl)-1-piperidinyl]carbonyl]-1H-pyrazol-5-yl]-
MF C23 H24 Cl N5 O2



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

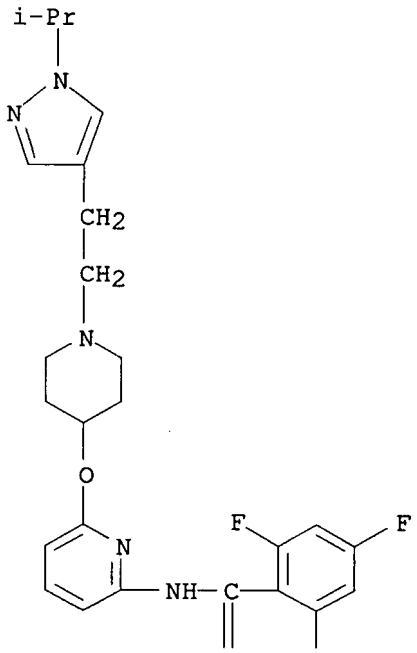
L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Carbamic acid, [2-[4-[1-(6-methoxy-3-pyridinyl)-3-(1-piperidinylcarbonyl)-1H-pyrazol-5-yl]phenoxy]ethyl]-, 1,1-dimethylethyl ester (9CI)
MF C28 H35 N5 O5



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Benzamide, 2,4,6-trifluoro-N-[6-[[1-[2-[1-(1-methylethyl)-1H-pyrazol-4-yl]ethyl]-4-piperidinyl]oxy]-2-pyridinyl]- (9CI)
MF C25 H28 F3 N5 O2
CI COM

PAGE 1-A



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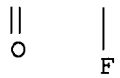
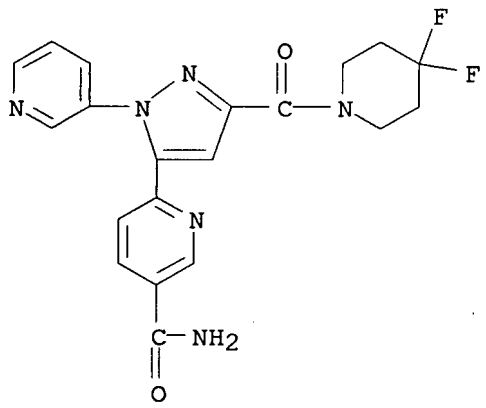


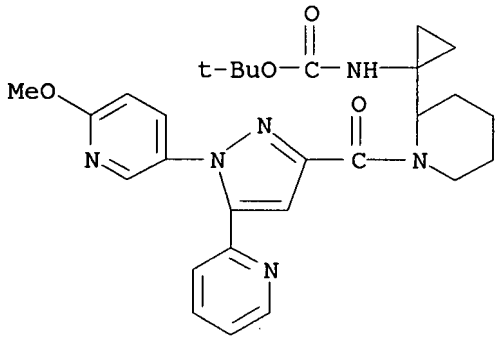
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 3-Pyridinecarboxamide, 6-[3-[(4,4-difluoro-1-piperidinyl)carbonyl]-1-(3-pyridinyl)-1H-pyrazol-5-yl]- (9CI)
MF C20 H18 F2 N6 O2



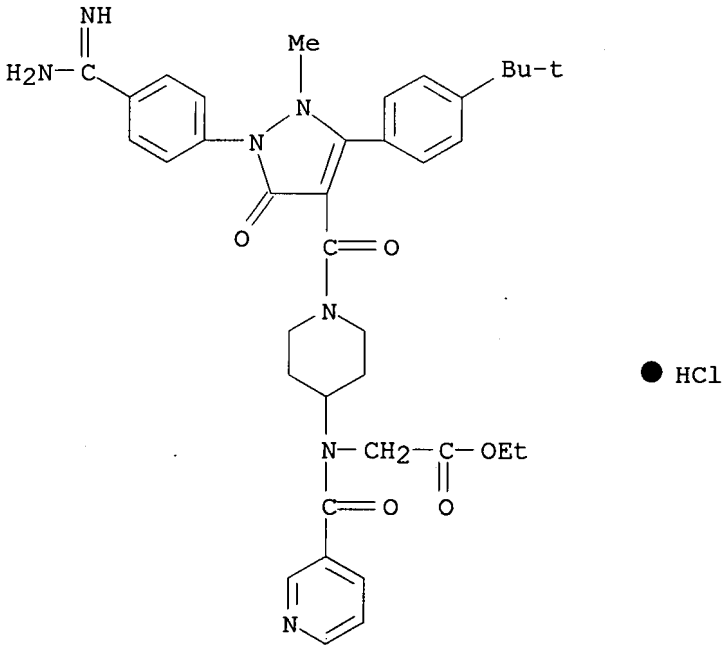
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Carbamic acid, [1-[1-[[1-(6-methoxy-3-pyridinyl)-5-(2-pyridinyl)-1H-pyrazol-3-yl]carbonyl]-2-piperidinyl]cyclopropyl]-, 1,1-dimethylethyl ester (9CI)
MF C28 H34 N6 O4



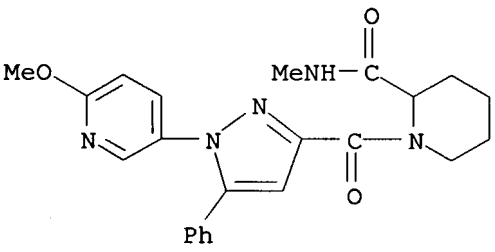
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Glycine, N-[1-[[2-[4-(aminoiminomethyl)phenyl]-5-[4-(1,1-dimethylethyl)phenyl]-2,3-dihydro-1-methyl-3-oxo-1H-pyrazol-4-yl]carbonyl]-4-piperidinyl]-N-(3-pyridinylcarbonyl)-, ethyl ester, monohydrochloride (9CI)
MF C37 H43 N7 O5 . Cl H



● HCl

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 2-Piperidinecarboxamide, 1-[[1-(6-methoxy-3-pyridinyl)-5-phenyl-1H-pyrazol-3-yl]carbonyl]-N-methyl- (9CI)
MF C23 H25 N5 O3



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Benzamide, 2-chloro-4-fluoro-N-[6-[methyl[1-[2-[1-(1-methylethyl)-1H-pyrazol-4-yl]ethyl]-4-piperidinyl]amino]-2-pyridinyl]-, monohydrochloride (9CI)
MF C26 H32 Cl F N6 O . Cl H

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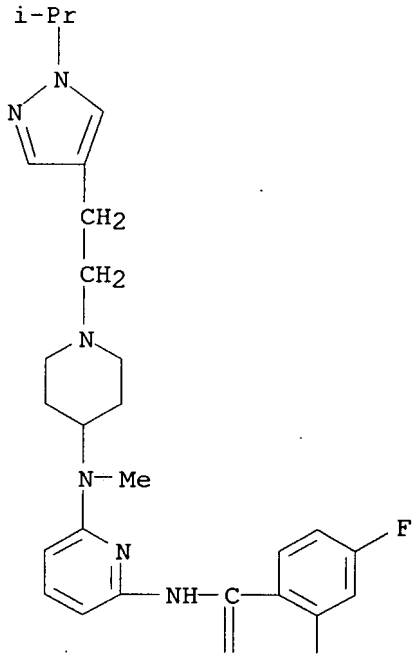

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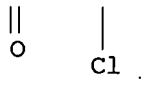
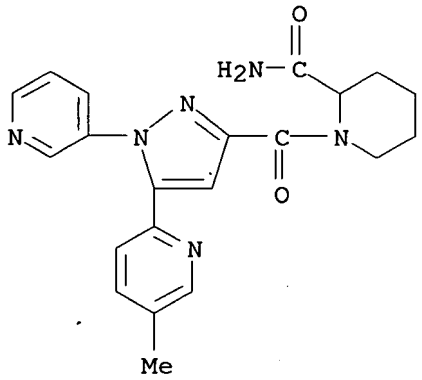


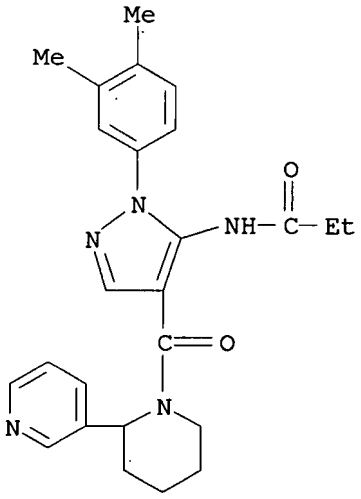
● HCl

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 2-Piperidinecarboxamide, 1-[[5-(5-methyl-2-pyridinyl)-1-(3-pyridinyl)-1H-pyrazol-3-yl]carbonyl]- (9CI)
MF C21 H22 N6 O2



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

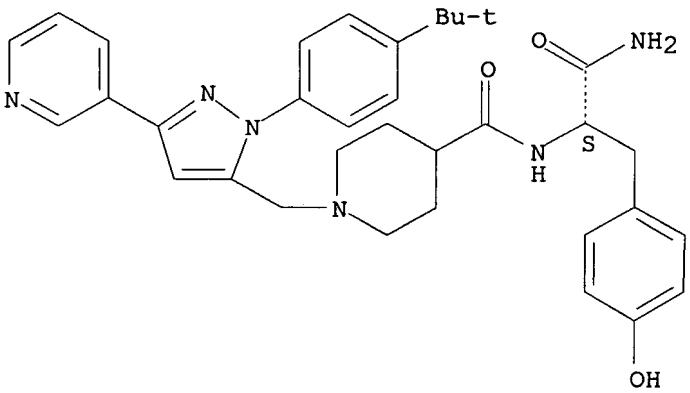
L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Propanamide, N-[1-(3,4-dimethylphenyl)-4-[[2-(3-pyridinyl)-1-piperidinyl]carbonyl]-1H-pyrazol-5-yl]-
MF C25 H29 N5 O2



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L8 47 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 4-Piperidinecarboxamide, N-[(1S)-2-amino-1-[(4-hydroxyphenyl)methyl]-2-oxoethyl]-1-[[1-[4-(1,1-dimethylethyl)phenyl]-3-(3-pyridinyl)-1H-pyrazol-5-yl]methyl]- (9CI)
MF C34 H40 N6 O3

Absolute stereochemistry.



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ALL ANSWERS HAVE BEEN SCANNED

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            20 C26 H33 F N6 O/MF
             6 C25 H28 F3 N5 O2 . CL H/MF
             2 C26 H32 CL F N6 O/MF
L9           5 L8 AND  ( C26 H32 CL F N6 O . CL H/MF OR  C25 H28 F3 N5 O2/MF
               OR  C26 H33 F N6 O/MF OR  C25 H28 F3 N5 O2 . CL H/MF OR   C26
              H32 CL F N6 O/MF )

=> fil caplus
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                                                           ENTRY    SESSION
FULL ESTIMATED COST                                       206.30     206.51

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=> s l9
L10          2 L9

=> d bib abs hitstr

L10  ANSWER 1 OF 2  CAPLUS  COPYRIGHT 2007 ACS on STN
AN   2005:346988  CAPLUS
DN   142:392299
TI   Preparation of aniline- and aminopyridine-derivatives as 5-HT1F receptor
     agonists
IN   Blanco-Pillado, Maria-Jesus; Cohen, Michael Philip; Filla, Sandra Ann;
     Hudziak, Kevin John; Kohlman, Daniel Timothy; Benesh, Dana Rae; Victor,
     Frantz; Xu, Yao-Chang; Ying, Bai-Ping; Zacherl, Deanna Piatt; Zhang, Deyi
PA   Eli Lilly and Company, USA
SO   PCT Int. Appl., 127 pp.
     CODEN: PIXXD2
DT   Patent
LA   English
FAN.CNT 1
     PATENT NO.          KIND   DATE        APPLICATION NO.          DATE
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PI WO 2005035499 A1 20050421 WO 2004-US25607 20040903
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RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
AU 2004280320 A1 20050421 AU 2004-280320 20040903
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CN 1849307 A 20061018 CN 2004-80026400 20040903
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JP 2007505105 T 20070308 JP 2006-526084 20040903
US 2006287363 A1 20061221 US 2006-569109 20060221
IN 2006KN00450 A 20070202 IN 2006-KN450 20060227
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WO 2004-US25607 W 20040903
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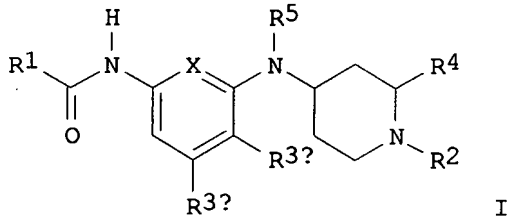

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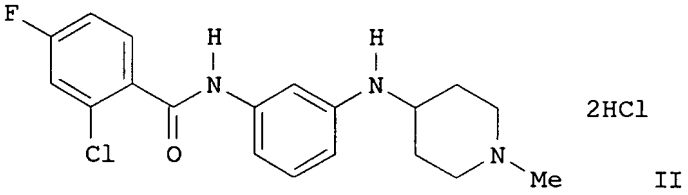

II

AB Title compds. I [X = -C(R3c)=, -N=; R1 = (un)substituted-alkyl, -cycloalkyl, -Ph, etc.; R2 = H, n-alkyl, cycloalkylalkyl with provisions; R3a, R3b, and, when X = -C(R3c)=, R3c independently = H, F, CH3 with provisions; R4 = H, alkyl; R5 = H, alkyl, cycloalkylcarbonyl with provisions] and their pharmaceutically acceptable salts, are prepared and disclosed as useful agonists for 5-HT1F receptor. Thus, e.g., II was prepared by reductive alkylation of 2-chloro-4-fluoro-N-(3-aminophenyl)benzamide (preparation given) with 1-methylpiperidin-4-one. The binding ability of I towards the 5-HT1F receptor was evaluated using radioligand binding assay and it revealed that selected compds. of the invention had a high affinity for the receptor, with exemplary Ki values

in the range of 600 nm or less. I as 5-HT1F receptor agonists should prove useful in the treatment of migraine.

IT 850082-67-6P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of aniline- and aminopyridine-derivs. as 5-HT1F receptor agonists)

RN 850082-67-6 CAPLUS

CN Benzamide, 2-chloro-4-fluoro-N-[6-[methyl[1-[2-[1-(1-methylethyl)-1H-pyrazol-4-yl]ethyl]-4-piperidinyl]amino]-2-pyridinyl]-, monohydrochloride (9CI) (CA INDEX NAME)

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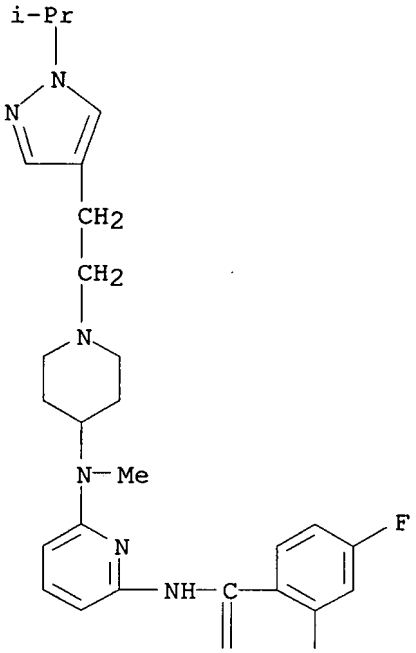

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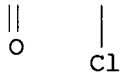

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RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d bib abs hitstr 2

L10 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2004:927173 CAPLUS
DN 141:395422

TI Preparation of N-[(piperidinyloxy)phenyl]-, N-[(piperidinyloxy)pyridinyl]-, N-[(piperidinylsulfanyl)phenyl]-, and N-[(piperidinylsulfanyl)pyridinyl] amides as 5-HT1F agonists for treatment of migraine

IN Blanco-Pillado, Maria-Jesus; Benesh, Dana Rae; Filla, Sandra Ann; Hudziak, Kevin John; Mathes, Brian Michael; Kohlman, Daniel Timothy; Ying, Bai-Ping; Zhang, Deyi; Xu, Yao-Chang

PA Eli Lilly and Company, USA

SO PCT Int. Appl., 186 pp.
CODEN: PIXXD2

DT Patent

LA English

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	CA 2518839	A1	20041104	CA 2004-2518839	20040414
	EP 1626958	A1	20060222	EP 2004-759769	20040414
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	BR 2004009211	A	20060328	BR 2004-9211	20040414
	CN 1777584	A	20060524	CN 2004-80010411	20040414
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OS MARPAT 141:395422

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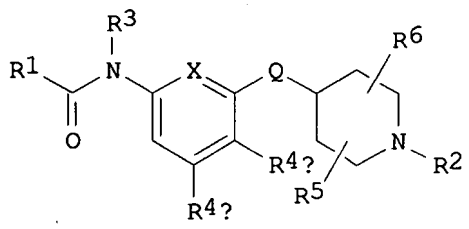

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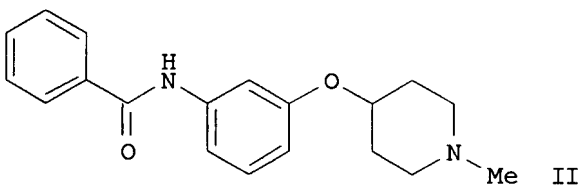

II

AB Title compds. I [wherein Q = O, S; X = CR4c, N; R1 = (un)substituted alkyl, cycloalkyl(alkyl), Ph, heterocyclyl; R2 = H, (fluoro)alkyl, cycloalkylalkyl, (un)substituted pyrazolyl(alkyl); R3 = H, alkyl; R4a, R4b, R4c = independently H, halo, (fluoro)alkyl; R5, R6 = independently H, (fluoro)alkyl; with the proviso that R6 = alkyl only when R5 ≠ H; and pharmaceutically acceptable acid addition salts thereof] were prepared by standard and solid phase combinatorial methods as 5-HT1F agonists. For example, amidation of [3-[(1-methylpiperidin-4-yl)oxy]phenyl]amine (preparation given) with benzoyl chloride afforded II (91%). In a radioligand binding assay using Ltk cells transfected with the human 5-HT1F receptor sequence, exemplified invention compds. exhibited high affinity for the receptor with Ki values of ≤ 150 nM. Thus, I and their pharmaceutical compns. are useful for activating 5-HT1F receptors, inhibiting neuronal protein extravasation, and treating or preventing migraine in mammals, especially humans (no data).

IT 790669-85-1P, 2,4,6-Trifluoro-N-[6-[[1-[2-(1-isopropyl-1H-pyrazol-4-yl)ethyl]piperidin-4-yl]oxy]pyridin-2-yl]benzamide 790669-86-2P, 2,4,6-Trifluoro-N-[6-[[1-[2-(1-isopropyl-1H-pyrazol-4-yl)ethyl]piperidin-4-yl]oxy]pyridin-2-yl]benzamide hydrochloride
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(5-HT1F agonist; preparation of piperidinyl-substituted amides as 5-HT1F agonists for treatment of migraine)

RN 790669-85-1 CAPLUS

CN Benzamide, 2,4,6-trifluoro-N-[6-[[1-[2-[1-(1-methylethyl)-1H-pyrazol-4-yl]ethyl]-4-piperidinyl]oxy]-2-pyridinyl]- (9CI) (CA INDEX NAME)

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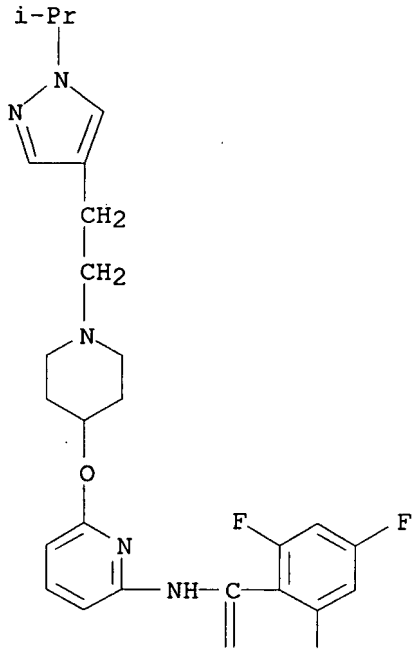

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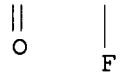

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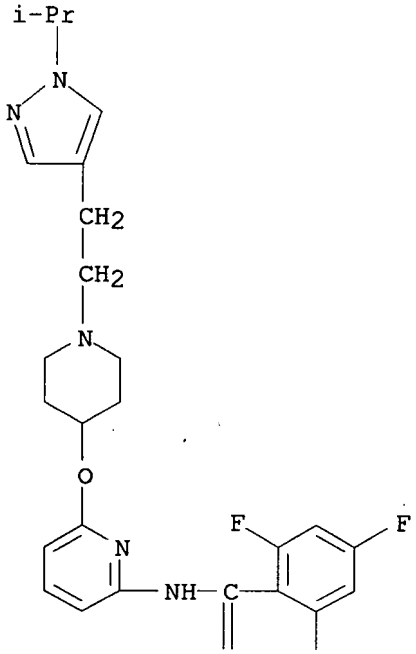
CN Benzamide, 2,4,6-trifluoro-N-[6-[[1-[2-[1-(1-methylethyl)-1H-pyrazol-4-yl]ethyl]-4-piperidinyl]oxy]-2-pyridinyl]-, monohydrochloride (9CI) (CA INDEX NAME)

PAGE 1-A



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● HCl

RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT